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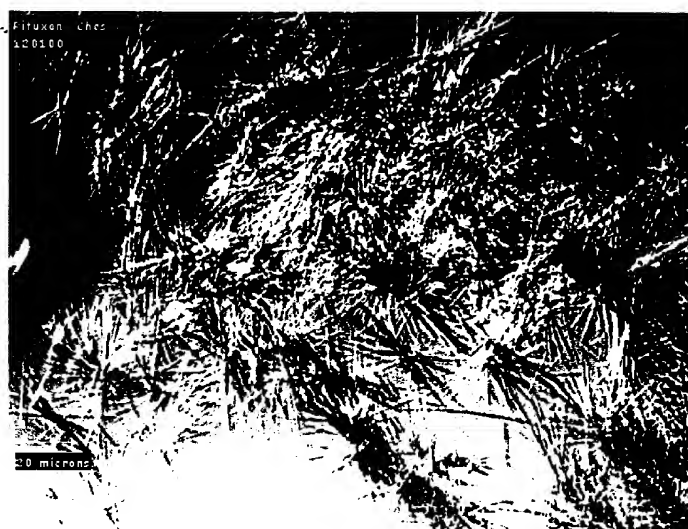
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(54) Title: CRYSTALS OF WHOLE ANTIBODIES AND FRAGMENTS THEREOF AND METHODS FOR MAKING AND USING THEM

### Rituximab Crystals



(57) Abstract: Methods are also provided for preparing stabilized formulations of whole antibody crystals or antibody fragment crystals using pharmaceutical ingredients or excipients and optionally encapsulating the crystals or crystal formulations in a polymeric carrier to produce compositions and using such protein crystals for biomedical applications, including delivery of therapeutic proteins and vaccines.



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CRYSTALS OF WHOLE ANTIBODIES AND FRAGMENTS  
THEREOF AND METHODS FOR MAKING AND USING THEM

This application claims the priority benefit  
of U.S. Provisional Application Serial No. 60/258,704,  
5 filed December 28, 2000, the content of which is  
incorporated by reference in its entirety.

TECHNICAL FIELD OF THE INVENTION

This invention relates to crystals of whole  
antibodies and fragments thereof, and formulations and  
10 compositions comprising such crystals. More  
particularly, methods are provided for the  
crystallization of high concentrations of whole  
antibodies, and fragments thereof, in large batches,  
and for the preparation of stabilized whole antibody  
15 crystals for use alone, or in dry or slurry  
formulations or compositions. This invention also  
relates to methods for stabilization, storage and  
delivery of biologically active whole antibody  
crystals.

20 The present invention further relates to  
methods using whole antibody crystals, antibody  
fragment crystals, or compositions or formulations  
comprising such crystals for biomedical applications,  
including biological delivery to humans and animals.

More particularly, highly concentrated whole antibody or antibody fragment crystal formulations or compositions are useful for delivery of large amounts of antibodies in a small volume to a subject, when and  
5 where they are needed. According to one embodiment of this invention, whole antibody crystals or antibody fragment crystals are used as a carrier-free delivery system which can slowly release active whole antibodies or fragments thereof, to a subject, where and when they  
10 are needed. According to an alternate embodiment of this invention, whole antibody crystals or antibody fragment crystals, or crystal formulations thereof, are encapsulated within a matrix comprising a polymeric carrier to form a composition.

15 Methods are also provided for preparing stabilized formulations of whole antibody crystals or antibody fragment crystals using pharmaceutical ingredients or excipients and optionally encapsulating the crystals or crystal formulations in a polymeric  
20 carrier to produce compositions and using such crystals for biomedical applications, including delivery of therapeutic proteins and vaccines.

#### BACKGROUND OF THE INVENTION

Antibodies, through their exquisite ability  
25 to specifically target a distinct antigen on an endogenous cell, bacteria, virus, or toxin, constitute powerful therapeutic agents characterized by limited side effects. Several antibodies introduced onto the market over the past few years have achieved  
30 astonishing success in treating a variety of diseases, including cancer and inflammatory, cardiovascular, respiratory, and infectious diseases. There are



currently approximately 480 launched and developmental antibody programs worldwide, 83% of which are located in the United States. Over 20% of all biopharmaceuticals currently being evaluated in  
5 clinical trials are antibodies, according to the Pharmaceutical Research Institute of America reports. The projected United States antibody market is anticipated to increase about ten-fold over the next decade, to \$10.1 billion in 2010 (*The Genesis Report:*  
10 *25+ Business Development & Innovation Opportunities in Monoclonal Antibodies-Emerging Opportunities in 2010*, The Genesis Group, Montclair, NJ). In contrast to such efforts in antibody development, techniques for their purification, stabilization or subsequent delivery are  
15 often limited.

It is imperative that the higher order three-dimensional architecture or tertiary structure of an antibody be preserved until such time that the individual antibody molecules are required to perform  
20 their unique function. To date, a limiting factor for the use of antibodies, particularly in therapeutic regimens, remains the sensitivity of antibody structure to chemical and physical denaturation encountered during delivery. Various approaches have been employed  
25 to overcome these barriers. However, these approaches often incur loss of protein activity or the additional expense of protein stabilizing carriers or formulations.

The stability of small molecule crystalline  
30 drugs is such that they can withstand extreme forces during the manufacturing process (see United States patent 5,510,118). Such forces are associated with milling nanoparticles of crystalline material of relatively insoluble drugs and include: shear stress,

turbulent flow, high impact collisions, cavitation and grinding. Small molecular crystalline compounds have been recognized as being much more stable toward chemical degradation than the corresponding amorphous solid [Pical, M.J., Lukes, A.L., Lang, J.E. and Gaines, J. Pharm. Sci. **67**:767 (1978)].

To date, those of skill in the art recognize that the greatly enhanced stability of the crystalline state observed for small molecules does not translate to biological macromolecules, such as whole antibodies [Pical, M.J. and Rigsbee, D.R., Pharm. Res. **14**:1379 (1997)]. For example, aqueous suspensions of crystalline insulin are only slightly more stable (to the degree of a factor of two) than corresponding suspensions of amorphous phase [Brange, J., Langkjaer, L., Havelund, S. and Volund, A., Pharm. Res. **9**:715 (1992)]. In the solid state, lyophilized amorphous insulin is more stable than lyophilized crystalline insulin under all conditions investigated so far [Pical, M.J. and Rigsbee, D.R., Pharm. Res. **14**:1379 (1997)]. However, using two model proteins, glucose oxidase and lipase, Shenoy et al. demonstrated that dry crystalline formulations can be significantly more stable than their amorphous counterparts [Shenoy, B. et al., Biotechnol Bioeng. **73**(5):358-69 (2001)].

Surprisingly, the present invention provides crystals of whole antibodies and crystals of single-chain Fv (scFv) antibody fragments or Fab antibody fragments (the "ab" stands for "antigen-binding") that are more stable than their soluble antibody or antibody fragment counterparts.

Despite recent progress in protein technology generally, two problems continue to limit the use of biological macromolecules in industry and medicine.

The first problem relates to molecular stability and sensitivity of higher order tertiary structures to chemical and physical denaturation during manufacturing and storage. Second, the field of biological delivery of therapeutic proteins requires that vehicles be provided which release native proteins, such as whole antibodies, at a rate that is consistent with the needs of the particular patient or disease process.

Although crystallization of whole antibodies has been a subject of significant interest for the last three decades, very few whole antibodies have ever been crystallized and, even then, solely in the context of structural studies [Harris L.J., Skaletsky, E., and McPherson, A., J. Mol. Biol. **275**:861-72 (1998); Harris L.J., Larson, S.B., Skaletsky, E., and McPherson, A., Immunological Reviews **163**:35-43 (1998)]. All of these crystals were obtained by vapor diffusion techniques, which yielded only a very small quantity of crystals for structural analysis. Such yields were far below those required for pharmaceutical, diagnostic or other commercial applications. Furthermore, such low yields were largely attributed to the difficulties in antibody crystallization due to their relatively large size, the presence of oligosaccharides on their surfaces, and the high degree of their segmental flexibility.

Fab antibody fragments have also been crystallized, but solely for use in X-ray crystallographic structural studies [See, e.g., Ito et al., Acta Crystallogr. D. Biol. Crystallogr. **57**:1700-02 (2001); Covaceuszach et al., Acta Crystallogr. D. Biol. Crystallogr. **57**:1307-09 (2001); Saul et al., Bioorg. Khim. **25**:247-52 (1999); Pichla et al., J. Struct. Biol. **119**:6-16 (1997); Maninder et al., J. Mol. Biol. **242**:706-08 (1994)].

The following table provides a general comparison between crystallization for X-ray crystallographic structural studies and large-scale crystallization according to this invention:

5	<b>PARAMETER</b>	<b>X-RAY CRYSTALLOGRAPHIC STUDIES</b>	<b>LARGE-SCALE CRYSTALLIZATION</b>
	Crystal size (longest dimension)	> 500 $\mu\text{m}$	0.1 - 100 $\mu\text{m}$
	Crystal quality	Very important	Less important
10	Growth rate	Not important	Important
	Yield	Not important	Very important
	Precipitate	Usually present	Rarely present

Crystallization of whole antibodies, or  
15 fragments thereof, on a large scale, a process allowing an alternative route of delivery for therapeutic antibodies, has never before been explored.

Antibodies, and fragments thereof, are increasingly employed in the pharmaceutical, diagnostic  
20 and research industries. There is a great need for alternative stabilization procedures, which are fast, inexpensive and Moreover, stabilization procedures are needed that do not involve the excessive use of  
25 whole antibodies.

The present invention seeks to overcome barriers to the widespread use of antibodies for therapeutic and other biomedical purposes by providing methods for crystallizing whole antibodies, and  
30 fragments thereof, on a large scale.

SUMMARY OF THE INVENTION

The present invention overcomes the above-described obstacles by employing the most stable form of an active, whole antibody or fragment thereof, the crystalline form. In one embodiment of this invention, crystals of a whole antibody, or fragment thereof are used as is or in formulations or compositions, for various biomedical applications. According to alternate embodiments of this invention, crystals of a whole antibody, or fragment thereof, or formulations or compositions comprising them, may be: (1) stabilized by adding ingredients or excipients to the crystals, or (2) encapsulated within a polymeric carrier to produce a composition that contains each crystal for delivery to a subject and subsequent release of active, whole antibodies. Any whole antibody or fragment thereof may be crystallized and/or stabilized in this manner, according to the methods of this invention.

Various aspects of this invention are particularly advantageous.

First, crystallinity of stored materials is very important, since large scale crystallization can be introduced as a purification step and/or concentration step in clinical manufacturing processes, such as those for manufacturing therapeutics and vaccines. Moreover, large scale crystallization can replace some of the purification steps in the manufacturing process. For example, whole antibody crystallization can streamline the production of antibody formulations and compositions, making the procedure more efficient and affordable.

Second, macromolecular interactions which occur in solution are prevented or severely reduced in the crystalline state, due to considerable reduction of all reaction rates. Thus, the crystalline state is uniquely suited to the storage of mixtures of whole antibodies, or fragments thereof.

Third, solid crystalline preparations may be easily reconstituted to generate ready to use parenteral preparations having very high antibody concentrations. Typically, for subcutaneous administration, injection volumes of 1.5 ml or less are well tolerated. Thus, for proteins that are dosed at 1 mg/kg on a weekly basis, a protein concentration of at least 50 mg/ml is required and 100-200 mg/ml is preferred. Such concentrations are difficult to achieve in liquid preparations, due to problems of aggregation and viscosity of the liquid samples. In contrast, they can be achieved in the crystalline preparations, or formulations or compositions thereof, according to this invention.

Fourth, whole antibody crystals, or crystals of antibody fragments, also constitute a particularly advantageous form for pharmaceutical dosage preparation. The crystals may be used as a basis for slow release *in vivo*. As those of skill in the art will appreciate, particle size is important for the dissolution of crystals and release of activity. Those skilled in the art will also appreciate the rate of antibody release to be more predictable if the crystals have substantially uniform particle size and do not contain amorphous precipitate. Thus, whole antibody crystals, or crystals of antibody fragments, may be advantageously used on implantable devices, such as those described in PCT patent application WO 96/40049.

Implant reservoirs are generally on the order of 25-250  
ul. With this volume restriction, a preparation of  
high concentration (greater than 10%) and a minimum  
amount of suspension vehicle is preferred. Whole  
5 antibody crystals, or crystals of antibody fragments,  
according to this invention may also be easily  
formulated in non-aqueous suspensions in such high  
concentrations.

Fifth, the use of whole antibody crystals, or  
10 crystals of antibody fragments, and formulations and  
compositions comprising them, for slow release of the  
antibody after delivery to the intended site,  
advantageously permits the effective biological half-  
life of the whole antibody or antibody fragment *in vivo*  
15 to be increased.

Sixth, another advantage of whole antibody  
crystals, or crystals of antibody fragments, is that  
certain variables can be manipulated to modulate the  
release of macromolecules over time. For example,  
20 crystal size, shape, formulation with excipients that  
effect dissolution, and encapsulation into a polymer  
matrix can all be manipulated to produce delivery  
vehicles for the antibodies.

The process of crystallization of whole  
25 antibodies serves not only as a powerful protein  
purification or stabilization tool but also affords the  
most concentrated protein form possible. Such effects  
have significant potential for delivery to the intended  
delivery site of a high dose of whole antibodies, or  
30 fragments thereof. Furthermore, by employing crystals  
or crystal formulations or compositions of whole  
antibodies or antibody fragments, for delivery to  
subjects, it is possible to carry out the controlled  
release of the whole antibodies or fragments thereof at

a rate that is consistent with the needs of the particular subject or disease process. As the rate of crystal dissolution depends on crystal morphology, crystal size and the presence of excipients, and the  
5 particular encapsulation technique or polymer preparation employed, crystalline whole antibodies or fragments thereof may also be used as a carrier-free slow release dosage form.

Whole antibodies or fragments thereof which  
10 are not stable when held in solution at ambient or elevated temperatures can nevertheless be successfully stored in dry crystalline form for long periods of time at such temperatures when they are crystallized according to the methods of this invention.

15                   BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts the morphology of Rituximab (Rituxan™) crystals prepared as described in Example 6. Rituximab crystals formed in needle clusters.

Figure 2 depicts the morphology of Infliximab  
20 (Remicade™) crystals prepared as described in Example 34. Infliximab crystals formed in rod-shaped clusters.

Figure 3 depicts the morphology of Rituximab (Rituxan™) crystals prepared as described in Example 28. Rituximab formed cube-shaped crystals.

25                   Figure 4 depicts the morphology of Rituximab (Rituxan™) crystals prepared as described in Example 26. Rituximab formed small needle-like crystals.

Figure 5 depicts the morphology of Trastuzumab (Herceptin™) crystals prepared as described  
30 in Example 31. Trastuzumab formed short needle-like crystals.



Figure 6 depicts the morphology of Trastuzumab (Herceptin™) crystals prepared as described in Example 32. Trastuzumab formed long needle-like crystals.

5           Figure 7 depicts the morphology of Infliximab (Remicade™) crystals prepared as described in Example 37. Trastuzumab formed star-shaped crystals.

          Figure 8 shows that crystallized Rituximab is capable of inducing a Direct Cytotoxicity response  
10   against the RAJI Lymphoma Cell Line. See Example 55.

          Figure 9 shows that crystallized Rituximab is capable of inducing Complement-Dependent Cytotoxicity against RAJI Lymphoma Cells. See Example 56.

          Figure 10 is a plot showing the results of an  
15   analysis of the stability of crystalline Rituximab in the presence of PEG, ethanol or a combination of PEG and ethanol. See Example 68.

          Figure 11 is a plot showing the results of an analysis of the stability of crystalline Trastuzumab  
20   (Herceptin™) in the presence of PEG, ethanol or a combination of PEG and ethanol. See Example 69.

          Figure 12 shows an SDS-PAGE gel of whole Rituximab antibody obtained by dissolving Rituximab crystals (as prepared on Example 1) that had been  
25   stored at room temperature for one month before being dissolved. See Example 64.

          Figure 13 is a chromatogram that depicts the results of treating crystalline Trastuzumab (Herceptin™) with acetone for three hours. The  
30   Trastuzumab remained whole and maintained its native structure. See Example 65.

          Figure 14 is a chromatogram that depicts the results of treating native (soluble) Trastuzumab (Herceptin™) with acetone for 20 minutes. The

native/soluble Trastuzumab precipitated after the acetone treatment, demonstrating a loss of the structural integrity of the native Trastuzumab. See Example 65.

5                Figure 15 is a plot that compares the rate at which crystalline Trastuzumab becomes bioavailable in the blood when administered intravenously (i.v.) with the rate of blood bioavailability when Trastuzumab is administered subcutaneously (s.c.).

10

#### DETAILED DESCRIPTION OF THE INVENTION

In order that the invention herein described may be more fully understood, the following detailed description is set forth. In the description, the  
15 following terms are employed:

Whole Antibody or Antibody Fragment -- a whole antibody or antibody fragment, e.g., a single-chain Fv fragment or Fab antibody fragment, according to this invention, is a functional antibody or antibody  
20 fragment, i.e., that is able to recognize and bind to its specific antigen *in vitro* or *in vivo*, and may initiate any subsequent actions associated with antibody-binding, e.g., Direct Cytotoxicity, Complement-Dependent Cytotoxicity (CDC), Antibody-  
25 Dependent Cytotoxicity (ADCC).

Amorphous solid -- a non-crystalline solid form of protein, sometimes referred to as "amorphous precipitate", which has no molecular lattice structure characteristic of the crystalline solid state.

30                Antibody -- a glycoprotein of approximate MW 150 kD, that is produced by the humoral arm of the immune system of vertebrates in response to the presence of foreign molecules in the body. Antibodies

are essential for the prevention and resolution of infection by microorganisms, e.g. parasites, bacteria and viruses. Antibodies perform this function by recognizing and binding, in a highly specific manner, protein (or, sometimes, other organic molecules including polysaccharides, glycoproteins, lipids, or nucleic acids) configurations called antigens (or epitopes), including those on invading microorganisms and their products. Antibodies bind their target antigens through highly specific interactions between hypervariable domains, called antigen-binding sites, on the antibody, and the epitope itself. Upon binding to the antigen, antibodies activate one or more of the many effector systems of the immune system that contribute to the neutralization, destruction and elimination of the infecting microorganism, or other antigen-containing entity, e.g. cancer cell.

Antibodies are also used for the treatment of cancer, inflammation, cardiovascular disease, and transplant rejection, by virtue of their specific binding and subsequent neutralization of the cellular targets, which are involved in disease states. For example, monoclonal antibody Infliximab binds to tumor necrosis factor and neutralizes its role in inflammation by blocking its interaction with cell surface receptor; while Rituximab targets malignant B lymphocytes by binding to their cell surface CD20 antigen.

A single antibody molecule has a structure composed of two identical heavy chains (each of approximate MW 50 kD) covalently bound to each other, and two identical light chains (each of approximate MW 25 kD), each covalently bound to one of the heavy chains. The four chains are arranged in a classic "Y"

motif. The bottom "leg" of the "Y" is called the Fc region ("c" stands for "crystallizable" or, alternatively, "complement-binding") and is used to anchor the antibody within cell membranes, and also to bind macrophage cells and activate complement. The two "arms" at the top of the "Y" are called Fab regions (the "ab" stands for "antigen-binding"). Each Fab region contains a constant region (at the juncture of the Fab and the Fc regions) and a variable region (which extends to the tip of the "Y"). Each variable region contains identical antigen-binding sites (at regions within the variable regions called "hypervariable" regions) at each tip of the "Y". Thus, each Fab region has one antigen-binding site, and the complete antibody molecule therefore has two antigen-binding sites (i.e., is "bivalent"). The two antigen-binding sites on a naturally occurring antibody are identical to each other, and therefore the antibody is specific for one antigen (i.e., is "monovalent"). A number of molecular fragments of antibody molecules have been isolated to date. These do not occur naturally, but are engineered from one or more complete antibody molecules. These fragments include Fab fragments (a single Fab that is isolated from a complete antibody by digestion with the enzyme papain), and  $F(ab')_2$  fragments (two Fabs covalently-bound to each other, produced by digesting the antibody with the enzyme pepsin). Fab fragments are monospecific, while  $F(ab')_2$  fragments are bispecific. Recently, a number of engineered antibody fragments have been introduced. These include double-stranded Fv (dsFv) fragments and single-chain Fv (scFv) fragments (the "v" stands for "variable" in both cases). A dsFv fragment consists of an Fab fragment minus the constant regions, i.e.,

consisting only of the variable regions of a heavy and light chain covalently bound to each other. A scFv fragment is a single polypeptide chain, consisting of the variable region of a heavy chain linked via a peptide linker to the variable region of a light chain. Classically, both dsFv and scFv fragments are monovalent (and thus mono-specific). However, two dsFv fragments or two scFv fragments can themselves be linked to form a bispecific fragment (which would be analogous to an  $F(ab')_2$  fragment without the constant regions). Furthermore, it is possible to link two dsFv fragments or scFv fragments with different antigen-binding sites (i.e., different specificities), to form a bi-specific fragment. Such fragments may be used as either research tools or therapeutic or diagnostic reagents.

There are five classes of antibodies (also called immunoglobulins) in humans: IgG, IgM, IgA, IgD, and IgE, each with its own unique characteristics and function. IgG, IgD, and IgE are all made up of one antibody molecule, while IgA can be made up of one, two or three such molecules and IgM consists of five. Furthermore, in humans, there are four subclasses of IgG (IgG1, IgG2, IgG3, or IgG4), and two subclasses each of IgM and IgA (1 and 2, respectively). For example, the monoclonal antibody Rituximab (Rituxan™) is an IgG1 antibody.

Though naturally occurring antibodies are derived from a single species, engineered antibodies and antibody fragments may be derived from more than one species of animal, i.e., may be chimeric. To date, mouse (murine)/human chimeric antibodies have been generated, though other species' combinations are possible. Chimeric antibodies have been further broken

down into two subtypes: chimeric and humanized. Chimeric murine/human antibodies contain approximately 75% human and 25% mouse amino acid sequences, respectively. The human sequences represent the  
5 constant regions of the antibody while the mouse sequences represent the variable regions (and thus contain the antigen-binding sites) of the antibody. The rationale for using such chimeras is to retain the antigen specificity of the mouse antibody but reduce  
10 the immunogenicity of the mouse antibody (a murine antibody would cause an immune response against it in species other than the mouse) and thus be able to employ the chimera in human therapies. Chimeric antibodies also include those which comprise CDR  
15 regions from different human antibodies. CDR regions, also called hypervariable regions, are sequences within the variable regions of antibody molecules that generate the antigen-binding sites. CDR regions are so-named because the binding site is complementary in  
20 shape and charge distribution to the epitope recognized on the antigen.

Alternatively, chimeric antibodies comprise framework regions from one antibody and CDR regions from another antibody. Chimeric antibodies also  
25 include those which comprise CDR regions from at least two different human antibodies. Humanized antibodies contain approximately 90% (or more) human amino acid sequences. The only murine sequences present are those for the hypervariable region (that are the actual  
30 antigen-binding sites contained within the variable region). Humanized antibodies have minimal mouse immunogenicity as compared with chimeric antibodies.

There are generally two types of antibodies that can be distinguished by their specificities:

polyclonal antibodies and monoclonal antibodies.

Polyclonal antibodies are those that are found as the immunoglobulin fraction of blood, and are essentially a polyclonal mixture of many different types of

- 5 antibodies specific for the different antigens the individual has been exposed to (i.e., they originate from many different clones of B lymphocytes (or B cells), the cell that produces antibodies).

- Monoclonal antibodies are antibodies of a  
10 single specificity, i.e., that are derived from a single clone of B lymphocytes (B cells). These antibodies have exquisite specificity for their target antigens and also can be produced in high amounts (i.e., high titres). They are useful as markers for  
15 specific antigens (e.g., cancer antigens), as diagnostic agents (e.g., in assays to detect viruses like HIV-1), and as therapeutic agents. Whole monoclonal antibodies are those that have a classic molecular structure that includes two complete heavy  
20 chains and two complete light chains. This is distinguished from antibody fragments, such as Fab, F(ab')<sub>2</sub>, Fc fragments, dsFv fragments, and scFv fragments.

- Traditionally, monoclonal antibodies have  
25 been produced by fusing the antibody-producing B cell with an immortal hybridoma cell to generate B cell hybridomas, which continually produce monoclonal antibodies in cell culture. Another method that is traditionally used to generate monoclonal antibodies  
30 involves the expression of the monoclonal antibodies in bacterial cell culture using phage-display technology. Currently, however, monoclonal antibodies may be produced *in vivo* in large quantities in genetically-modified animals, such as cows and goats (Genzyme

Transgenics), pigs and rabbits (Medarex, PPL Therapeutics), and chickens (Tranxenogen), and in plants, such as tobacco and corn (Epicyte, Integrated Protein Technologies, Meristem CropTech, and others).

5 For example, large amounts of monoclonal antibodies can be found in the milk of genetically-modified goats (Genzyme Transgenics). Antibodies from all such sources may be crystallized according to this invention. Furthermore, as a result of transgenics,

10 mice have been modified to contain and express the entire human B cell genome (which encodes human antibodies). Therefore, such transgenic mice (Abgenix) are a source of human antibodies for crystallization according to this invention. It should be noted that

15 glycosylation is specific to the animal that is producing the antibodies. For example, human antibodies from sources other than humans will have subtly different glycosylation profiles. Therefore, the whole antibodies or single-chain Fv antibody

20 fragments or Fab antibody fragments of this invention may display modified glycosylation or be deglycosylated. Antibodies which may be crystallized according to this invention also include derivatized antibodies. Such antibodies include those derivatized

25 with polyethylene glycol or at least one carbohydrate moiety or least one methyl or ethyl group. Clinically relevant antibodies may also be classified according to the therapeutic area in which they are to be employed. Such antibodies include, for example, those for

30 treating cancers (e.g., pancreatic cancer), inflammatory diseases (e.g., autoimmune diseases, arthritis), cardiovascular diseases (e.g., strokes), infectious disease (e.g., HIV/AIDS), respiratory diseases (e.g., asthma), tissue transplantation



rejection and organ transplantation rejection. Such antibodies also include antibodies for radioimmunotherapy. Antibodies which may be crystallized according to the present invention  
5 include, for example, Abciximab, Palivizumab, Murumonab-CD3, Gemtuzumab, Trastuzumab, Basiliximab, Daclizumab, Etanercept and Ibritumomab tiuxetan.

Antibody activity release rate -- the quantity of whole antibody, single-chain Fv antibody  
10 fragment or Fab antibody fragment dissolved per unit time.

Antigen -- any substance or material that is specifically recognized and bound by an antibody. Antigens are typically small pieces of proteins  
15 (peptides) found on the surfaces of cells or invading microorganisms. Antibodies are thought to specifically recognize antigens as small as four amino acids in length, and the substitution of only one amino acid can abolish antibody recognition of the particular antigen  
20 for which it is specific.

Antigenicity -- the ability of an antigen to be specifically recognized and bound by an antibody. An antigen is said to be in its antigenic conformation when it can be specifically recognized and bound by the  
25 antibody specific for the antigen. This is different from immunogenicity, which is the ability of an antigen to elicit the production of antibodies specific for the antigen.

Anti-idiotypic antibody -- antibodies having  
30 specificity for the antigen-binding sites of other antibody molecules. Anti-idiotypic antibodies are generated in the following manner: an antigen elicits the production of antibodies (called Ab-1 or idiotypes) that are specific for that antigen. These antibodies

(idiotypes) are then used as immunogens themselves to elicit a second generation of antibodies that are specific for Ab-1. These second generation antibodies (Ab-2) are called anti-idiotypic antibodies (or anti-  
5 idiotypes), and either mimic, or are closely related to, the initial antigen used to generate Ab-1. Such reactions also occur naturally *in vivo*, in response to antigenic stimulation, and by means of these antibody-antibody interactions, the immune system is able to, in  
10 essence, interact with itself. It has been postulated that by exploiting this capability, anti-idiotypic antibodies can be used to prevent certain infections, and treat some kinds of cancers and various immune and autoimmune diseases.

15       Antibody half-life -- for antibodies *in vivo*, the time in which a given amount of whole antibody, a single-chain Fv antibody fragment or Fab antibody fragment, are reduced to 50% of its initial concentration. IgG typically has a half-life of about  
20 21 days (though IgG3 has a half-life of only 7 days), while IgM, A, D, and E have typical half-lives of 10 days, 6 days, 3 days, and 2 days, respectively.

25       Antibody loading -- the antibody content of formulations or compositions, as calculated as a percentage by weight of antibody, a single-chain Fv antibody fragment or Fab antibody fragment, relative to the weight of the dry preparation. A typical range of antibody loading is from 1-80%.

30       Antibody release -- the release of active protein from a polymeric carrier, as controlled by one or more of the following factors: (1) degradation of the polymer matrix; (2) rate of crystal dissolution within the polymer matrix; (3) diffusion of dissolved protein through the polymer matrix; (4) protein

loading; and (5) diffusion of biological medium into the antibody crystal/polymer matrix.

Aqueous-organic solvent mixture -- a mixture comprising n% organic solvent, where n is between 1 and 5 99 and m% aqueous, where m is 100-n.

Bioavailability -- the degree to which a substance, e.g., an active antibody or antibody fragment, administered *in vivo*, becomes available to the tissue to which the substance is targeted.

10 According to this invention, bioavailability also refers to the degree to which a whole antibody, or fragment thereof, that has been administered *in vivo* as a crystal or a composition or formulation thereof, becomes available in the blood. According to this

15 invention, bioavailability also refers to the ability of the substance, e.g., an active antibody or antibody fragment, to perform a function, e.g., direct cytotoxicity, at the target tissue once the substance has been delivered. Bioavailability may be measured in

20 a number of ways, e.g., as the concentration of the substance, e.g., an active antibody or antibody fragment, measured as a function of time in the bloodstream.

Biocompatible polymers -- polymers that are

25 non-antigenic (when not used as an adjuvant), non-carcinogenic, non-toxic and which are not otherwise inherently incompatible with living organisms. Examples include: poly (acrylic acid), poly (cyanoacrylates), poly (amino acids), poly

30 (anhydrides), poly (depsipeptide), poly (esters) such as poly (lactic acid) or PLA, poly (lactic-co-glycolic acid) or PLGA, poly ( $\beta$ -hydroxybutyrate), poly (caprolactone) and poly (dioxanone); poly (ethylene glycol), poly ((hydroxypropyl)methacrylamide, poly

[(organo)phosphazene], poly (ortho esters), poly (vinyl alcohol), poly (vinylpyrrolidone), maleic anhydride-alkyl vinyl ether copolymers, pluronic polyols, albumin, alginate, cellulose and cellulose derivatives, collagen, fibrin, gelatin, hyaluronic acid, oligosaccharides, glycaminoglycans, sulfated polysaccharides, blends and copolymers thereof.

Biodegradable polymers -- polymers that degrade by hydrolysis or solubilization. Degradation can be heterogenous -- occurring primarily at the particle surface, or homogenous -- degrading evenly throughout the polymer matrix.

Biological macromolecule -- biological polymers such as proteins, deoxyribonucleic acids (DNA) and ribonucleic acids (RNA). For the purposes of this application, biological macromolecules are also referred to as macromolecules.

Composition -- crystals of whole antibodies or crystals of single-chain Fv antibody fragments or Fab antibody fragments, or formulations thereof, which have been encapsulated within a polymeric carrier to form coated particles.

Controlled dissolution -- dissolution of a crystal of an whole antibody or single-chain Fv antibody fragment or Fab antibody fragment, or a formulation or composition comprising such crystals, or release of the crystalline constituent of said crystal or formulation or composition that is controlled by a factor selected from the group consisting of the following: the surface area of said crystal; the size of said crystal; the shape of said crystal; the concentration of excipient component of the formulation or composition; the number and nature of excipient components of the formulation or composition; the

molecular weight of the excipient components of the formulation or composition; the nature of the polymeric carriers, and combinations thereof.

Co-polymer -- a polymer made with more than  
5 one monomer species.

Crystal -- Crystals are one form of the solid state of matter, which is distinct from a second form - the amorphous solid state, which exists essentially as an unorganized, heterogeneous solid. Crystals are  
10 regular three-dimensional arrays of atoms, ions, molecules (e.g., proteins such as antibodies), or molecular assemblies (e.g., antigen/antibody complexes). Crystals are lattice arrays of building blocks called asymmetric units (which consist of the  
15 substance to be crystallized) that are arranged according to well-defined symmetries into unit cells that are repeated in three-dimensions. See Giegé, R. and Ducruix, A. Barrett, *Crystallization of Nucleic Acids and Proteins, a Practical Approach*, 2nd ed., pp.  
20 1-16, Oxford University Press, New York, New York, (1999).

Dissolution of crystals -- dissolving a crystal of a whole antibody or fragment thereof in order to recover soluble antibodies or antibody  
25 fragments.

Drying of Crystals of Whole Antibodies or Single-chain Fv Antibody Fragments or Fab Antibody Fragments -- removal of water, organic solvent or liquid polymer by means including drying with N<sub>2</sub>, air or  
30 inert gases, vacuum oven drying, lyophilization, washing with a volatile organic solvent followed by evaporation of the solvent, evaporation in a fume hood, tray drying, fluid bed drying, spray drying, vacuum drying, or roller drying. Typically, drying is

achieved when the crystals become a free flowing powder. Drying may be carried out by passing a stream of gas over wet crystals. The gas may be selected from the group consisting of: nitrogen, argon, helium,  
5 carbon dioxide, air or combinations thereof.

Effective amount -- an amount of a crystal of an whole antibody or a crystal of a single-chain Fv antibody fragment or Fab antibody fragment or crystal formulation or composition of this invention which is  
10 effective to treat, immunize, boost, protect, repair or detoxify the subject or area to which it is administered over some period of time.

Emulsifier -- a surface active agent which reduces interfacial tension between polymer coated  
15 crystals and a solution.

Formulation - a combination of crystals of an whole antibody, or a combination of crystals of single-chain Fv antibody fragment or crystals of an Fab antibody fragment, and one or more ingredients or  
20 excipients, including sugars and biocompatible polymers. Examples of excipients are described in the *Handbook of Pharmaceutical Excipients*, published jointly by the American Pharmaceutical Association and the Pharmaceutical Society of Great Britain. For the  
25 purposes of this application, "formulations" include "crystal formulations." Furthermore, "formulations" include "whole antibody crystal formulations" and "single-chain Fv antibody fragment crystal formulations" and "Fab antibody crystal formulations".

30 Glycoprotein -- a protein or peptide covalently linked to a carbohydrate. The carbohydrate may be monomeric or composed of oligosaccharides.

Homo-polymer -- a polymer made with a single monomer species.

Immunotherapeutic -- an antibody or single-chain Fv antibody fragment or Fab antibody fragment is immunotherapeutic when it has the activity of inducing protective immunity to a tumor cell, virus, or bacteria  
5 or stimulating the immune system to reduce or eliminate said tumor cell, virus or bacteria.

Ingredients -- any excipient or excipients, including pharmaceutical ingredients or excipients. Excipients include, for example, the following:

10 **Acidifying agents**

acetic acid, glacial acetic acid, citric acid, fumaric acid, hydrochloric acid, diluted hydrochloric acid, malic acid, nitric acid, phosphoric acid, diluted phosphoric acid, sulfuric acid, tartaric  
15 acid

**Aerosol propellants**

butane, dichlorodifluoromethane, dichlorotetrafluoroethane, isobutane, propane, trichloromonofluoromethane

20 **Air displacements**

carbon dioxide, nitrogen

**Alcohol denaturants**

denatonium benzoate, methyl isobutyl ketone, sucrose octacetate

25 **Alkalizing agents**

strong ammonia solution, ammonium carbonate, diethanolamine, diisopropanolamine, potassium hydroxide, sodium bicarbonate, sodium borate, sodium carbonate, sodium hydroxide, trolamine

5 **Anticaking agents** (see *glidant*)

**Antifoaming agents**

dimethicone, simethicone

**Antimicrobial preservatives**

10 benzalkonium chloride, benzalkonium chloride solution, benzelthonium chloride, benzoic acid, benzyl alcohol, butylparaben, cetylpyridinium chloride, chlorobutanol, chlorocresol, cresol, dehydroacetic acid, ethylparaben, methylparaben, methylparaben sodium, phenol, phenylethyl alcohol, phenylmercuric acetate, phenylmercuric nitrate, potassium benzoate, 15 potassium sorbate, propylparaben, propylparaben sodium, sodium benzoate, sodium dehydroacetate, sodium propionate, sorbic acid, thimerosal, thymol

**Antioxidants**

20 ascorbic acid, acorbyl palmitate, butylated hydroxyanisole, butylated hydroxytoluene, hypophosphorous acid, monothioglycerol, propyl gallate, sodium formaldehyde sulfoxylate, sodium metabisulfite, sodium thiosulfate, sulfur dioxide, tocopherol, 25 tocopherols excipient



**Buffering agents**

acetic acid, ammonium carbonate, ammonium phosphate, boric acid, citric acid, lactic acid, phosphoric acid, potassium citrate, potassium  
5 metaphosphate, potassium phosphate monobasic, sodium acetate, sodium citrate, sodium lactate solution, dibasic sodium phosphate, monobasic sodium phosphate, histidine

**Capsule lubricants** (see *tablet and capsule lubricant*)

**10 Chelating agents**

edetate disodium, ethylenediaminetetraacetic acid and salts, edetic acid

**Coating agents**

sodium carboxymethylcellulose, cellulose  
15 acetate, cellulose acetate phthalate, ethylcellulose, gelatin, pharmaceutical glaze, hydroxypropyl cellulose, hydroxypropyl methylcellulose, hydroxypropyl methylcellulose phthalate, methacrylic acid copolymer, methylcellulose, polyethylene glycol, polyvinyl acetate  
20 phthalate, shellac, sucrose, titanium dioxide, carnauba wax, microcrystalline wax, zein

**Colors**

caramel, red, yellow, black or blends, ferric oxide

**Complexing agents**

ethylenediaminetetraacetic acid and salts (EDTA), edetic acid, gentisic acid ethanolamide, oxyquinoline sulfate

**5 Desiccants**

calcium chloride, calcium sulfate, silicon dioxide

**Emulsifying and/or solubilizing agents**

acacia, cholesterol, diethanolamine  
10 (adjunct), glyceryl monostearate, lanolin alcohols, lecithin, mono- and di-glycerides, monoethanolamine (adjunct), oleic acid (adjunct), oleyl alcohol (stabilizer), poloxamer, polyoxyethylene 50 stearate, polyoxyl 35 castor oil, polyoxyl 40 hydrogenated castor  
15 oil, polyoxyl 10 oleyl ether, polyoxyl 20 cetostearyl ether, polyoxyl 40 stearate, polysorbate 20, polysorbate 40, polysorbate 60, polysorbate 80, propylene glycol diacetate, propylene glycol monostearate, sodium lauryl sulfate, sodium stearate,  
20 sorbitan monolaurate, sorbitan monooleate, sorbitan monopalmitate, sorbitan monostearate, stearic acid, trolamine, emulsifying wax

**Filtering aids**

powdered cellulose, purified siliceous earth

**25 Flavors and perfumes**

anethole, benzaldehyde, ethyl vanillin,  
menthol, methyl salicylate, monosodium glutamate,  
orange flower oil, peppermint, peppermint oil,  
peppermint spirit, rose oil, stronger rose water,  
5 thymol, tolu balsam tincture, vanilla, vanilla  
tincture, vanillin

**Glidant and/or anticaking agents**

calcium silicate, magnesium silicate,  
10 colloidal silicon dioxide, talc

**Humectants**

glycerin, hexylene glycol, propylene glycol,  
sorbitol

**Ointment bases**

15 lanolin, anhydrous lanolin, hydrophilic  
ointment, white ointment, yellow ointment, polyethylene  
glycol ointment, petrolatum, hydrophilic petrolatum,  
white petrolatum, rose water ointment, squalane

**Plasticizers**

20 castor oil, lanolin, mineral oil, petrolatum,  
benzyl benyl formate, chlorobutanol, diethyl phthalate,  
sorbitol, diacetylated monoglycerides, diethyl  
phthalate, glycerin, glycerol, mono- and di-acetylated  
monoglycerides, polyethylene glycol, propylene glycol,  
25 triacetin, triethyl citrate, ethanol

**Polymer membranes**

cellulose acetate

#### **Solvents**

acetone, alcohol, diluted alcohol, amylene hydrate, benzyl benzoate, butyl alcohol, carbon  
5 tetrachloride, chloroform, corn oil, cottonseed oil, ethyl acetate, glycerin, hexylene glycol, isopropyl alcohol, methyl alcohol, methylene chloride, methyl isobutyl ketone, mineral oil, peanut oil, polyethylene glycol, propylene carbonate, propylene glycol, sesame  
10 oil, water for injection, sterile water for injection, sterile water for irrigation, purified water

#### **Sorbents**

powdered cellulose, charcoal, purified siliceous earth  
15 **Carbon dioxide sorbents**

barium hydroxide lime, soda lime

#### **Stiffening agents**

hydrogenated castor oil, cetostearyl alcohol, cetyl alcohol, cetyl esters wax, hard fat, paraffin,  
20 polyethylene excipient, stearyl alcohol, emulsifying wax, white wax, yellow wax

#### **Suppository bases**

cocoa butter, hard fat, polyethylene glycol

**Suspending and/or viscosity-increasing agents**

acacia, agar, alginic acid, aluminum monostearate, bentonite, purified bentonite, magma bentonite, carbomer 934p, carboxymethylcellulose  
5 calcium, carboxymethylcellulose sodium, carboxymethylcellulose sodium 12, carrageenan, microcrystalline and carboxymethylcellulose sodium cellulose, dextrin, gelatin, guar gum, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropyl  
10 methylcellulose, magnesium aluminum silicate, methylcellulose, pectin, polyethylene oxide, polyvinyl alcohol, povidone, propylene glycol alginate, silicon dioxide, colloidal silicon dioxide, sodium alginate, tragacanth, xanthan gum

**15 Sweetening agents**

aspartame, dextrates, dextrose, excipient dextrose, fructose, mannitol, saccharin, calcium saccharin, sodium saccharin, sorbitol, solution sorbitol, sucrose, compressible sugar, confectioner's  
20 sugar, syrup

**Tablet binders**

acacia, alginic acid, sodium carboxymethylcellulose, microcrystalline cellulose, dextrin, ethylcellulose, gelatin, liquid glucose, guar  
25 gum, hydroxypropyl methylcellulose, methylcellulose, polyethylene oxide, povidone, pregelatinized starch, syrup

**Tablet and/or capsule diluents**

calcium carbonate, dibasic calcium phosphate,  
tribasic calcium phosphate, calcium sulfate,  
microcrystalline cellulose, powdered cellulose,  
dextrans, dextrin, dextrose excipient, fructose,  
5 kaolin, lactose, mannitol, sorbitol, starch,  
pregelatinized starch, sucrose, compressible sugar,  
confectioner's sugar

**Tablet disintegrants**

alginate acid, microcrystalline cellulose,  
10 croscarmellose sodium, croscopolone, polacrillin  
potassium, sodium starch glycolate, starch,  
pregelatinized starch

**Tablet and/or capsule lubricants**

calcium stearate, glyceryl behenate,  
15 magnesium stearate, light mineral oil, polyethylene  
glycol, sodium stearyl fumarate, stearic acid, purified  
stearic acid, talc, hydrogenated vegetable oil, zinc  
stearate

**Tonicity agent**

20 dextrose, glycerin, mannitol, potassium  
chloride, sodium chloride

**Vehicle: flavored and/or sweetened**

aromatic elixir, compound benzaldehyde  
elixir, iso-alcoholic elixir, peppermint water,  
25 sorbitol solution, syrup, tolu balsam syrup

**Vehicle: oleaginous**

almond oil, corn oil, cottonseed oil, ethyl  
oleate, isopropyl myristate, isopropyl palmitate,  
mineral oil, light mineral oil, myristyl alcohol,  
5 octyldodecanol, olive oil, peanut oil, persic oil,  
sesame oil, soybean oil, squalane

**Vehicle: solid carrier**

sugar spheres

10 **Vehicle: sterile**

Bacteriostatic water for injection,  
bacteriostatic sodium chloride injection

**Viscosity-increasing** (see *suspending agent*)**Water repelling agent**

15 cyclomethicone, dimethicone, simethicone

**Wetting and/or solubilizing agent**

benzalkonium chloride, benzethonium chloride,  
cetylpyridinium chloride, docusate sodium, nonoxynol 9,  
nonoxynol 10, octoxynol 9, poloxamer, polyoxyl 35  
20 castor oil, polyoxyl 40, hydrogenated castor oil,  
polyoxyl 50 stearate, polyoxyl 10 oleyl ether, polyoxyl  
20, cetostearyl ether, polyoxyl 40 stearate,  
polysorbate 20, polysorbate 40, polysorbate 60,  
polysorbate 80, sodium lauryl sulfate, sorbitan

monolaureate, sorbitan monooleate, sorbitan  
monopalmitate, sorbitan monostearate, tyloxapol

Preferred ingredients or excipients include:

- 1) salts of amino acids such as glycine, arginine,
- 5 aspartic acid, glutamic acid, lysine, asparagine, glutamine, proline, histidine; 2) carbohydrates, e.g. monosaccharides such as glucose, fructose, galactose, mannose, arabinose, xylose, ribose; 3) disaccharides, such as lactose, trehalose, maltose, sucrose; 4)
- 10 polysaccharides, such as maltodextrins, dextrans, starch, glycogen; 5) alditols, such as mannitol, xylitol, lactitol, sorbitol; 6) glucuronic acid, galacturonic acid; 7) cyclodextrins, such as methyl cyclodextrin, hydroxypropyl- $\beta$ -cyclodextrin and alike 8)
- 15 inorganic salts, such as sodium chloride, potassium chloride, magnesium chloride, phosphates of sodium and potassium, boric acid ammonium carbonate and ammonium phosphate; 9) organic salts, such as acetates, citrate, ascorbate, lactate; 10) emulsifying or solubilizing
- 20 agents like acacia, diethanolamine, glyceryl monostearate, lecithin, monoethanolamine, oleic acid, oleyl alcohol, poloxamer, polysorbates, sodium lauryl sulfate, stearic acid, sorbitan monolaurate, sorbitan monostearate, and other sorbitan derivatives, polyoxyl
- 25 derivatives, wax, polyoxyethylene derivatives, sorbitan derivatives; and 11) viscosity increasing reagents like, agar, alginic acid and its salts, guar gum, pectin, polyvinyl alcohol, polyethylene oxide, cellulose and its derivatives propylene carbonate,
- 30 polyethylene glycol, hexylene glycol and tyloxapol. A further preferred group of excipients or ingredients includes sucrose, trehalose, lactose, sorbitol, lactitol, inositol, salts of sodium and potassium such



as acetate, phosphates, citrates, borate, glycine, arginine, polyethylene oxide, polyvinyl alcohol, polyethylene glycol, hexylene glycol, methoxy polyethylene glycol, gelatin, hydroxypropyl- $\beta$ -  
5 cyclodextrin.

Insoluble and stable form -- a form of crystal of an whole antibody or a single-chain Fv antibody fragment crystal or an Fab antibody fragment crystal which is insoluble in aqueous solvents, organic  
10 solvents or aqueous-organic solvent mixtures and which displays greater stability than the soluble form of the counterpart antibody or single-chain Fv antibody fragment or Fab antibody fragment. According to one embodiment of this invention, the phrase "insoluble and  
15 stable form" may denote a form of crystals which is insoluble in dry preparations but soluble in wet preparations. In any embodiment, the whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment may be  
20 active in insoluble form. And in one embodiment, the whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment may be active in insoluble form, then dissolve or are removed or digested once their function is  
25 complete. According to another embodiment of this invention, crystals of whole antibodies or fragments thereof may be crosslinked for added stability. According to another embodiment of this invention, metal ions, e.g.,  $\text{Ca}^{++}$ , may be added to crystals of  
30 whole antibodies or fragments thereof, rendering the crystals more insoluble and more stable.

Label -- incorporation of a label to a crystal of an whole antibody or of a single-chain Fv antibody fragment or of an Fab antibody fragment.

Labels may be selected from the group consisting of radiolabels, enzyme labels, toxins, magnetic agents or drug conjugates.

Liquid polymer -- pure liquid phase synthetic  
5 polymers, such as poly-ethylene glycol (PEG), in the absence of aqueous or organic solvents.

Loss of shelf stability -- the loss of specific activity and/or changes in secondary structure of a crystalline whole antibody or of a crystalline  
10 single-chain antibody Fv fragment or of a crystalline Fab antibody fragment as compared with the soluble (i.e., non-crystallized, native) antibody or single-chain Fv antibody fragment or Fab antibody fragment counterpart over time, when incubated under  
15 corresponding conditions.

Loss of stability -- the loss of specific activity and/or changes in secondary structure of a crystalline whole antibody or of a crystalline single-chain antibody Fv fragment or of a crystalline Fab  
20 antibody fragment as compared with the soluble (i.e., non-crystallized) antibody or single-chain Fv antibody fragment or Fab antibody fragment counterpart over time, while in solution under corresponding conditions.

Macromolecules -- proteins, glycoproteins, peptides, therapeutic proteins, DNA or RNA molecules,  
25 polysaccharides, lipoproteins, lipopolysaccharides.

Method of Administration -- crystals of whole antibodies or single-chain Fv antibody fragment crystals or Fab antibody fragment crystals, or crystal  
30 formulations or compositions thereof, may be appropriate for a variety of modes of administration. These may include oral and parenteral administration. Examples of parenteral administration according to this invention include, but are not limited to,

subcutaneous, intravenous, transdermal, intramuscular, pulmonary inhalation, intralesional, topical administration, needle injection, dry powder inhalation, skin electroporation, aerosol delivery, and  
5 needle-free injection technologies, including needle-free sub-cutaneous administration.

Microspheres -- encapsulated crystalline material which is spherical or roughly or nearly spherical, and has a diameter between about 1 nm and  
10 about 1 mm.

Microparticulates -- encapsulated crystalline material which has a diameter between about 1 nm and about 1 mm, but has no defined shape.

Mother Liquor -- the buffer used for  
15 crystallization of macromolecules, e.g., proteins, nucleic acids.

Needle-free drug delivery devices and jet injections -- delivery of a substance into the body of a mammal, or other suitable recipient, which does not  
20 involve using a sharp needle for injection. This may be a needle-free device which delivers the substance in a pressure-mediated manner. Examples of commercially-available needle-free injection devices or systems that can be used to administer crystals or crystal  
25 formulations or compositions of whole antibodies or antibody fragments according to this invention include, *inter alia*, Intraject™ (Weston Medical, Ltd.), Biojector2000® (Bioject, Inc.), MadaJet™ (MADA Medical Products, Inc.), and J-Tip® (National Medical Products, Inc.),  
30 LectraJet™ (DCI, Inc.), Mesoflash® (also called Isojet™) (Prolitec), VACCI JET Electrique™ (ENDOS Pharma), and a two-stage fluid medicament jet injector (Avant Drug Delivery Systems, Inc.).

Organic solvents -- any solvent of non-aqueous origin, including liquid polymers and mixtures thereof. Organic solvents suitable for the present invention include: acetone, methyl alcohol, methyl  
5 isobutyl ketone, chloroform, 1-propanol, isopropanol, 2-propanol, acetonitrile, 1-butanol, 2-butanol, ethyl alcohol, cyclohexane, N-methylpyrrolidinone (NMP), dioxane, ethyl acetate, dimethylformamide, dichloroethane, hexane, isooctane, methylene chloride,  
10 tert-butyl alcohol, toluene, carbon tetrachloride, or combinations thereof.

Pharmaceutically effective amount -- an amount of a crystal of an whole antibody or of a single-chain Fv antibody fragment or of an Fab antibody  
15 fragment, or crystal formulation or composition thereof, which is effective to treat a condition in an living organism to whom it is administered over some period of time.

Plasticizing -- use of a plasticizer, e.g.,  
20 lanolin, ethanol, to make a formulation comprising a whole antibody crystal or antibody fragment crystal in a solution that becomes viscous after it is injected subcutaneously, forming a matrix. The resulting high viscosity matrix is adhesive, biodegradable and  
25 biocompatible. The antibody or antibody fragment is then released in a controlled manner from the matrix.

Polyethylene glycol (PEG) size -- The size of the PEG moieties used according to this invention (e.g., *inter alia*, PEG 200, PEG 400, PEG 10,000, PEG  
30 80,000) refers to the chain length, i.e., number of ethylene glycol residues in the PEG chain. For example, PEG 200 has 200 ethylene glycol residues in the PEG polymer, PEG 80,000 has 80,000 ethylene glycol residues in the PEG polymer, etc.

Polymer -- a large molecule built up by the repetition of small, simple chemical units. The repeating units may be linear or branched to form interconnected networks. The repeat unit is usually  
5 equivalent or nearly equivalent to the monomer.

Polymeric carriers -- polymers used for encapsulation of whole antibody crystals or crystals of single-chain Fv antibody fragments or crystals of Fab antibody fragments for delivery of such whole  
10 antibodies or antibody fragments, including biological delivery. Such polymers include biocompatible and biodegradable polymers. The polymeric carrier may be a single polymer type or it may be composed of a mixture of polymer types. Polymers useful as the polymeric  
15 carrier, include for example, poly (acrylic acid), poly (cyanoacrylates), poly (amino acids), poly (anhydrides), poly (depsipeptide), poly (esters) such as poly (lactic acid) or PLA, poly (lactic-co-glycolic acid) or PLGA, poly ( $\beta$ -hydroxybutyrate), poly  
20 (caprolactone) and poly (dioxanone); poly (ethylene glycol), poly ((hydroxypropyl)methacrylamide, poly [(organo)phosphazene], poly (ortho esters), poly (vinyl alcohol), poly (vinylpyrrolidone), maleic anhydride-alkyl vinyl ether copolymers, pluronic polyols,  
25 albumin, natural and synthetic polypeptides, alginate, cellulose and cellulose derivatives, collagen, fibrin, gelatin, hyaluronic acid, oligosaccharides, glycaminoglycans, sulfated polysaccharides, modified starches such as amylose starch, amylopectin starch,  
30 hydroxyethyl starch, methacrylate starch, and other starches, and any conventional material that will encapsulate protein crystals.

Prophylactically effective amount -- an amount of a crystal of a whole antibody or single-chain

Fv antibody fragment crystal or Fab antibody fragment crystal, or crystal formulation or composition thereof, which is effective to prevent a condition in an living organism to whom it is administered over some period of  
5 time.

Protein -- a complex polymer containing carbon, hydrogen, oxygen, nitrogen and usually sulfur and composed of chains of amino acids connected by peptide linkages. The molecular weight range for  
10 proteins includes peptides of 1000 Daltons to glycoproteins of 600 to 1000 kilodaltons.

Protein delivery system -- method or means for administering one or more of a protein, such as an antibody crystal, single-chain Fv antibody fragment  
15 crystal, Fab antibody fragment crystal, or formulation or composition comprising such crystals, to a biological entity.

Radiolabel -- incorporation of a radiolabel to a protein, such as a crystal of an whole antibody or  
20 of a single-chain Fv antibody fragment or of an Fab antibody fragment. In situations where the radiolabel has a short half-life, as with  $^{131}\text{I}$  or  $^{90}\text{Y}$ , the radiolabel can also be therapeutic, e.g., used in radioimmunotherapies against cancers. Various methods  
25 of labeling polypeptides and glycoproteins are known in the art and may be used. Examples of labels include, but are not limited to, the following radioisotopes or radionucleotides:  $^3\text{H}$ ,  $^{14}\text{C}$ ,  $^{15}\text{N}$ ,  $^{35}\text{S}$ ,  $^{90}\text{Y}$ ,  $^{99}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{125}\text{I}$ , and  $^{131}\text{I}$ .

30 Reconstitution -- dissolution of whole antibody crystals or crystals of a single-chain Fv antibody fragment or of an Fab antibody fragment, or formulations or compositions comprising such crystals, in an appropriate buffer or pharmaceutical preparation.

Room Temperature -- for purposes of this invention, it will be understood by those of skill in the art that room temperature can be any temperature from about 20°C to about 26°C.

5           Stabilization -- the process of preventing the loss of specific activity and/or changes in secondary structure of a crystalline whole antibody or of a crystalline single-chain antibody Fv fragment or of a crystalline Fab antibody fragment as compared with  
10 the soluble antibody or single-chain Fv antibody fragment counterpart, or Fab antibody fragment counterpart, by preparing formulations or compositions of antibody or single-chain Fv antibody fragment or Fab antibody fragment crystals, with excipients or  
15 ingredients, including polymeric carriers.

Therapeutic antibody or single-chain Fv antibody fragment or Fab antibody fragments -- a crystal of a whole antibody or single-chain Fv antibody fragment or Fab antibody fragment, or crystal  
20 composition or formulation thereof, according to this invention, which is administered to a living organism to treat a given illness or symptom thereof.

Vaccine antibody or single-chain Fv antibody fragment or Fab antibody fragments -- an antibody or  
25 single-chain Fv antibody fragment or Fab antibody fragment that is elicited by (1) a native antigen, e.g., an antigen found on a pathogenic agent such as a virus, parasite, bacteria or tumor cell, or found on a tumor, or (2) an allergen. The protein activity of  
30 such vaccine antibodies or single-chain Fv antibody fragments or Fab antibody fragments is the induction of protective immune responses specific for a pathogenic agent, tumor, or allergen, or other antigen.

Crystallinity of Whole Antibodies and Fragments Thereof  
the Advantages Thereof

Crystallinity of macromolecules, such as whole antibodies, or fragments thereof, is of great value for their storage and delivery in vivo. However, few techniques exist for the preparation of large quantities of such crystalline macromolecules which are stable outside of the mother liquor. Crystals of proteins, such as whole antibodies and fragments thereof, must be handled with considerable care, since they are extremely fragile and contain a high proportion of solvent. It is well known in x-ray crystallography that the diffraction patterns from macromolecular crystals quickly degenerate upon dehydration in air. Normally, a crystal is carefully separated from its mother liquor and inserted into a capillary tube. The tube is sealed from the air using dental wax or silicone grease, along with a small amount of mother liquor inside to maintain hydration [McPherson, A., Preparation and Analysis of Protein Crystals, Robert E. Krieger Publishing, Malabar, p. 214 (1989)]. Another technique is to collect data from macromolecular crystals at cryogenic temperatures. The crystals are prepared and then rapidly cooled to prevent ice lattice formation in the aqueous medium. Instead of ice, a rigid glass forms, encasing the crystal with little damage. Crystals are then maintained at 100 °K to prevent crystal disintegrations [Rodgers, D.W., in Methods in Enzymology (Eds., Carter, C.W. and Sweet, R.M.) Academic Press, v.276, p. 183 (1997)]. While this technique allows one to maintain crystals outside of their mother liquor, it cannot be used at temperatures higher than 100 °K.



In principle, dried crystals can be prepared by lyophilization. However, this technique involves rapid cooling of the material and can be applied only to freeze stable products. The aqueous solution  
5 containing a crystalline whole antibody or a crystalline single-chain antibody Fv fragment or a crystalline Fab antibody fragment is first frozen to between -40 and -50°C. Then, the ice is removed under vacuum. Ice formation is usually destructive to the  
10 protein crystal lattice, yielding a mixture of crystals and amorphous precipitate.

It is desirable to produce whole antibodies, in the crystalline state, that are pure and stable under storage conditions at ambient temperatures. Such  
15 crystals constitute a particularly advantageous form for dosage preparations of therapeutics and vaccines. The present invention advantageously provides formulations and compositions of crystals of whole antibodies. The present invention also provides  
20 formulations and compositions for storage of crystals of whole antibodies as either solid particles or dispersed in a non-aqueous solvent. Furthermore, the invention may be applied to the storage of a single type of biologically active whole antibody or a mixture  
25 of different types of whole antibodies that do not interact with each other.

In another embodiment, this invention provides a method for crystallizing single-chain Fv (scFv) fragments of antibodies, and using such crystals  
30 in various biomedical applications. Such scFv fragments are constructed by linking the variable region of an antibody heavy chain to a variable region of an antibody light chain through the use of a linker peptide. Due to their small size, scFv fragments allow

tissue penetration more readily than do whole antibodies, and therefore may have valuable therapeutic applications for particular indications. It should be understood that crystals, crystal formulations or  
5 crystal compositions containing scFv fragments can be generated and utilized in the same manner applicable to crystals of whole antibodies, in the various embodiments of this invention.

In another embodiment, this invention  
10 provides a method for crystallizing Fab fragments of antibodies, and using such crystals in various biomedical applications. Such Fab fragments are generated by digesting a complete antibody with the enzyme papain, to yield antibody fragment molecules  
15 with one antigen binding site, as described above. Alternatively, Fab fragments may be generated by using genetic engineering technology. Due to their smaller size, Fab fragments allow tissue penetration more readily than do whole antibodies, and therefore may  
20 have particularly valuable therapeutic applications for particular indications. It should be understood that crystals, crystal formulations or crystal compositions containing Fab fragments can be generated and utilized in the same manner applicable to crystals of whole  
25 antibodies, in the various embodiments of this invention.

This invention allows crystallization of, and use of crystals of, all of the immunoglobulin classes IgG, IgM, IgA, IgD, IgE, and serum IgA (sIgA) as well  
30 as the subclasses IgG1, IgG2, IgG3 and IgG4, IgM1 and IgM2, and IgA1 and IgA2, as well as scFv fragments and Fab antibody fragments, from all the immunoglobulin classes and subclasses.

In another embodiment, this invention provides a method for rendering biologically active crystals of whole antibodies suitable for storage in suspensions comprising replacing the mother liquor with  
5 a non-aqueous solvent. In yet another embodiment, the crystalline slurry can be rendered solid by spinning out the first solvent and washing the remaining crystalline solid using a second organic solvent to remove water, followed by evaporation of the non-  
10 aqueous solvent.

Non-aqueous slurries of crystalline whole antibodies or scFv fragments or Fab fragments are especially useful for subcutaneous delivery, and intramuscular delivery, while solid preparations are  
15 ideally suited for pulmonary administration. As will be appreciated by those of skill in the art, pulmonary delivery is particularly useful for biological macromolecules which are difficult to deliver by other routes of administration.

20 Crystals of whole antibodies and crystals of single-chain Fv antibody fragments and crystals of Fab antibody fragments according to this invention are useful in diagnostic methods and kits. For example, such crystals may be used in a kit for diagnosing the  
25 presence a target antigen in a sample from a patient or another specimen. Such a kit may comprise a container and, optionally instructions for use. The crystals in the kit may be labelled with a detectable label. Methods for detecting a target antigen in a sample,  
30 such as a blood, tumor, cell, or tissue sample, may be carried out by mixing the sample with crystals of whole antibodies or crystals of single-chain Fv antibody fragments or Fab antibody fragments according to this invention and determining whether the sample binds to

the antibody or fragment. The crystals used in such methods may be labelled with a detectable label.

Alternatively, crystals of whole antibodies or crystals of single-chain Fv antibody fragments and  
5 crystals of Fab antibody fragments according to this invention are useful in chromatography and purification methods, such as affinity chromatography. For example, affinity matrix purification of a protein may be carried out by:

- 10 (a) mixing with a binding buffer crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, wherein such antibody or antibody fragment has affinity for the protein to be purified;
- 15 (b) adding a protein solution containing the protein to be purified to the crystal/buffer mixture;
- (c) incubating the entire mixture for a time and at a temperature sufficient to permit binding of the protein to the antibody or antibody fragment;
- 20 (d) washing the mixture with a wash buffer;
- and
- (e) eluting the protein with an elution buffer.

#### Stability of Encapsulated Crystals of Whole Antibodies

25 Those of skill in the art will appreciate that protein stability is one of the most important obstacles to the successful preparation of polymer microparticulate delivery systems that control the release of proteins. The stability of crystalline  
30 proteins, such as crystals of a whole antibody, or crystals of antibody fragments, encapsulated in polymeric carriers may be challenged at three separate stages: 1) manufacture of the antibody crystal

composition, 2) antibody release from the resulting composition and 3) in vivo stability after the antibody release. During preparation of microparticles or microspheres containing soluble or amorphous proteins, the use of organic solvents and lyophilization are especially detrimental to protein stability. Subsequently, released proteins are susceptible to moisture-induced aggregation, thus resulting in permanent inactivation.

10 In order to achieve high protein stability during preparation of whole antibody crystals, crystals of antibody fragments, or formulations and compositions according to the present invention, it is necessary to restrict the mobility of individual whole antibody molecules -- a result best achieved in the crystalline solid state. For the purpose of this application, solid state may be divided into two categories: amorphous and crystalline. The three-dimensional long-range order that normally exists in a crystalline material does not exist in the amorphous state. Furthermore, the position of molecules relative to one another is more random in the amorphous or liquid states, relative to the highly ordered crystalline state. Thus, amorphous proteins, including antibodies, may be less stable than their crystalline counterparts.

#### Maintaining Crystallinity

In order to use antibody crystals or antibody fragment crystals as the antibody source for preparing antibody formulations and compositions according to the present invention, the problem of protein crystal dissolution outside the crystallization solution ("mother liquor") had to be overcome. In order to maintain protein crystallinity, and hence stability, in

the production of the crystals of whole antibodies, or crystals of antibody fragments, and formulations and compositions of this invention, several approaches may be used:

- 5 1. Crystals remain in the mother liquor in the course of producing antibody crystals encapsulated with polymeric carriers. Many compounds used in protein crystallization, such as salts, PEG and organic solvents, are compatible with polymer  
10 processing conditions.
2. Kinetics of dissolution. The rate of crystal dissolution outside the mother liquor depends on conditions, such as pH, temperature, presence of metal ions, such as Zn, Cu and Ca and  
15 concentration of precipitants. By varying these conditions, one can slow down the dissolution of crystals for several hours. At the same time, the process of microparticulate formation is very fast and normally takes seconds to minutes to complete.
- 20 3. Dried antibody crystals. The mother liquor can be removed by filtration and the remaining crystalline paste can be dried by air, under vacuum, by washing with water miscible organic solvents and/or by lyophilization or spray drying.
- 25 4. The crystal size and shape can be manipulated and controlled in the course of crystallization. Thus, a range of crystal morphologies, each having different dissolution kinetics and subsequently different sustained release profiles compared to  
30 amorphous proteins, is available.

5. Method of making a crystal formulation by exchanging the mother liquor to a pharmaceutically-acceptable solvent or solution to form a Subcutaneous Vehicle for controlled delivery *in vivo*: The mother liquor can be removed by centrifugation and the remaining crystalline material can be suspended in a pharmaceutically acceptable solvent (e.g., ethanol) for subcutaneous injections. The crystalline material may also be suspended in sucrose acetate isobutyrate (SAIB) or poly (lactic-co-glycolic acid) (PLGA) in N-methylpyrrolidinone (NMP), where it forms a gel under the skin once it comes into contact with aqueous body fluids. The gel then facilitates the controlled release of the antibody or fragment thereof.

#### Administration and Biological Delivery

To date, therapeutic proteins, such as antibodies, have generally been administered by frequent injection or infusion, due to their characteristic negligible oral bioavailability and short plasma life. Crystals of whole antibodies, or crystals of antibody fragments, as well as crystal formulations and compositions containing them, according to the present invention, (which include microparticulate-based sustained release systems for whole antibodies), advantageously permit improved patient compliance and convenience. Furthermore, because of increased bioavailability and increased stability of proteins in the crystalline state, more stable blood levels of the administered antibodies or antibody fragments can be achieved, potentially with lower dosages. Also, the slow and constant release

capabilities afforded by the present invention advantageously permit reduced dosages, due to more efficient delivery of active antibody. Significant cost savings may be achieved by using the crystallized  
5 antibodies and antibody formulations and compositions described herein.

The antibody crystals, crystal formulations and compositions of the present invention enhance preservation of the native biologically active tertiary  
10 structure of the whole antibodies and create a reservoir which can slowly release active whole antibodies, or fragments thereof, to a subject where and when they are needed. The biologically active whole antibody, or fragment thereof, is subsequently  
15 released in a controlled manner over a period of time, as determined by the particular encapsulation technique, polymer constitution, crystal morphology, crystal size, crystal solubility, and the presence and nature of any excipients used. The crystals, crystal  
20 formulations and compositions of this invention may be reconstituted with a diluent for the parenteral administration of biologically active whole antibodies or antibody fragments.

Formulations and compositions comprising  
25 crystals of a whole antibody, or fragments thereof, in polymeric delivery carriers according to this invention may also comprise any conventional carrier or adjuvant used in vaccines, pharmaceuticals, personal care formulations and compositions, veterinary preparations,  
30 or oral enzyme supplementation. These carriers and adjuvants include, for example, Freund's adjuvant, ion exchangers, alumina, aluminum stearate, lecithin, buffer substances, such as phosphates, glycine, sorbic acid, potassium sorbate, partial glyceride mixtures of



saturated vegetable fatty acids, water, salts or electrolytes, such as protamine sulfate, disodium hydrogen phosphate, sodium chloride, zinc salts, colloidal silica, magnesium, trisilicate, cellulose-  
5 based substances and polyethylene glycol. Adjuvants for topical or gel base forms may include, for example, sodium carboxymethylcellulose, polyacrylates, polyoxyethylene-polyoxypropylene-block polymers, polyethylene glycol and wood wax alcohols.

10           According to one embodiment of this invention, crystals of an whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment may be combined with any conventional materials used for controlled release  
15 administration, including pharmaceutical controlled release administration. Such materials include, for example, coatings, shells and films, such as enteric coatings and polymer coatings and films.

Formulations or compositions comprising  
20 crystals of an whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment may be delivered to humans, animals, or plants at the desired site of delivery according to this invention. Such delivery may include the use of  
25 devices, such as implant-capable devices, or may involve other microparticulate protein delivery systems.

In one embodiment of this invention, crystals of an whole antibody or of a single-chain Fv antibody  
30 fragment or of an Fab antibody fragment have a longest dimension between about 0.01  $\mu\text{m}$  and about 500  $\mu\text{m}$ , alternatively between about 0.1  $\mu\text{m}$  and about 200  $\mu\text{m}$ . The most preferred embodiment is that the whole antibody crystals, or crystals of an antibody fragment,

are between about 1  $\mu\text{m}$  and about 100  $\mu\text{m}$  in their longest dimension. Such crystals may have a shape selected from the group consisting of: needles, needle clusters, disks, cubes, rods, quasi-crystals, spheres, plates, such as hexagons and squares, rhomboids, bipyramids and prisms, and others.

In one embodiment of this invention, formulations or compositions have a whole antibody concentration greater than about 0.1 mg/ml or a single-chain Fv antibody fragment concentration greater than about 0.1 mg/ml, or an Fab antibody fragment concentration greater than about 0.1 mg/ml. Alternatively, formulations or compositions have a whole antibody concentration greater than about 1 mg/ml or a single-chain Fv antibody fragment concentration greater than about 1 mg/ml, or an Fab antibody fragment concentration greater than about 1 mg/ml. Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 10 mg/ml or a single-chain Fv antibody fragment concentration greater than about 10 mg/ml, or an Fab antibody fragment concentration greater than about 10 mg/ml. Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 20 mg/ml or a single-chain Fv antibody fragment concentration greater than about 20 mg/ml, or an Fab antibody fragment concentration greater than about 20 mg/ml. Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 50 mg/ml or a single-chain Fv antibody fragment concentration greater than about 50 mg/ml, or an Fab antibody fragment concentration greater than about 50 mg/ml.

Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 100 mg/ml or a single-chain Fv antibody fragment concentration greater than about 100  
5 mg/ml, or an Fab antibody fragment concentration greater than about 100 mg/ml. Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 120 mg/ml or a single-chain Fv antibody fragment  
10 concentration greater than about 120 mg/ml, or an Fab antibody fragment concentration greater than about 120 mg/ml. Alternatively, formulations or compositions of the present invention have a whole antibody concentration greater than about 200 mg/ml or a single-  
15 chain.Fv antibody fragment concentration greater than about 200 mg/ml, or an Fab antibody fragment concentration greater than about 200 mg/ml.

According to this invention, any individual, including humans, animals and plants, may be treated in  
20 a pharmaceutically acceptable manner with a pharmaceutically effective amount of crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising  
25 such crystals, for a period of time sufficient to treat a condition in the individual to whom they are administered over some period of time. Alternatively, individuals may receive a prophylactically effective amount of whole antibody crystals or crystals of a  
30 single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, of this invention which is effective to prevent a condition in the individual to whom they are administered over some period of time.

Crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, may be administered alone, as  
5 part of a pharmaceutical, personal care or veterinary preparation, or as part of a prophylactic preparation, with or without adjuvant. They may be administered by parenteral or oral routes. For example, they may be administered by oral, pulmonary, nasal, aural, anal,  
10 dermal, ocular, intravenous, intramuscular, intraarterial, intraperitoneal, mucosal, sublingual, subcutaneous, transdermal, topical or intracranial routes, or into the buccal cavity. In either pharmaceutical, personal care or veterinary  
15 applications, crystals of whole antibodies or fragments thereof, or crystal formulations or compositions thereof may be topically administered to any epithelial surface. Such epithelial surfaces include oral, ocular, aural, anal and nasal surfaces, which may be  
20 treated, protected, repaired or detoxified by application of crystals of a whole antibody, crystals of a single-chain Fv antibody fragment, or crystals of an Fab antibody fragment, or crystal formulations or compositions thereof.

25           Pharmaceutical, veterinary or prophylactic preparations comprising crystals of a whole antibody or crystals of a single-chain Fv antibody fragment, or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, according to  
30 this invention may also be selected from the group consisting of tablets, liposomes, granules, spheres, microparticles, microspheres and capsules.

For such uses, as well as other uses according to this invention, crystals of a whole

antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, may be prepared in tablet form. Such tablets

5 constitute a liquid-free, dust-free form for storage of whole antibody crystals, crystals of antibody fragments, or crystal formulations or compositions which are easily handled and retain acceptable levels of activity or potency.

10 Alternatively, crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, may be in a variety of conventional forms employed for  
15 administration to provide reactive whole antibodies or single-chain Fv antibody fragments or Fab antibody fragments at the site where needed. These include, for example, solid, semi-solid and liquid dosage forms, such as liquid solutions or suspensions, slurries,  
20 gels, creams, balms, emulsions, lotions, powders, sprays, foams, pastes, ointments, salves, balms and drops.

Crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab  
25 antibody fragment, or formulations or compositions comprising such crystals, according to this invention may also comprise any conventional carrier or adjuvant used in pharmaceuticals, personal care preparations or veterinary preparations. These carriers and adjuvants  
30 include, for example, Freund's adjuvant, ion exchangers, alumina, aluminum stearate, lecithin, buffer substances, such as phosphates, glycine, sorbic acid, potassium sorbate, partial glyceride mixtures of saturated vegetable fatty acids, water, salts or

electrolytes, such as protamine sulfate, disodium hydrogen phosphate, sodium chloride, zinc salts, colloidal silica, magnesium, trisilicate, cellulose-based substances and polyethylene glycol. Adjuvants  
5 for topical or gel base forms may include, for example, sodium carboxymethylcellulose, polyacrylates, polyoxyethylene-polyoxypropylene-block polymers, polyethylene glycol and wood wax alcohols.

The most effective mode of administration and  
10 dosage regimen of crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, of this invention will depend on the effect desired, previous  
15 therapy, if any, the individual's health status, the status of the condition itself, the response to the whole antibody crystals or single-chain Fv antibody fragment crystals or Fab antibody fragment crystals, or crystal formulations or compositions thereof, and the  
20 judgment of the treating physician or clinician. The whole antibody crystals, single-chain Fv antibody fragment crystals, Fab antibody fragment crystals, or crystal formulations or compositions thereof, may be administered in any dosage form acceptable for  
25 pharmaceuticals, immunotherapy, or veterinary preparations, at one time or over a series of treatments.

The amount of crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or  
30 crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, which provides a single dosage will vary depending upon the particular mode of administration, the specific crystal preparation, formulation or composition, dose level and

dose frequency. A typical preparation will contain between about 0.01% and about 99%, preferably between about 1% and about 50%, of whole antibody crystals or crystals of a single-chain Fv antibody fragment or  
5 crystals of an Fab antibody fragment, or formulations or compositions thereof (w/w). Alternatively, a preparation will contain between about 0.01% and about 80% whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab  
10 antibody fragment, or formulations or compositions thereof (w/w), preferably between about 1% and about 50%, antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations or compositions thereof  
15 (w/w).

Upon improvement of the individual's condition, a maintenance dose of crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or  
20 formulations or compositions comprising such crystals, may be administered, if necessary. Subsequently, the dosage or frequency of administration, or both, may be reduced as a function of the symptoms, to a level at which the improved condition is retained. When the  
25 condition has been alleviated to the desired level, treatment should cease. Individuals may, however, require intermittent treatment on a long-term basis upon any recurrence of the condition or symptoms thereof.

30 The crystallized whole antibodies, single-chain Fv antibody fragments and Fab antibody fragments, and compositions and formulations thereof, according to this invention, may be used to treat a wide variety of human and other diseases, infections and disorders

including, *inter alia*, any human diseases which can be treated with antibodies, alone or in combination with other drugs, or in complex or conjugated with other chemical substances, e.g., toxins or radionucleotides.

5 Diseases, infections and disorders which may be treated or diagnosed using the crystallized whole antibodies, scFv antibody fragments and Fab antibody fragments, and compositions and formulations thereof, according to this invention, include, *inter alia*: AIDS/HIV infection

10 or related conditions; autoimmune disorders like rheumatoid arthritis, systemic lupus erythematosus, idiopathic thrombocytopenic purpura; blood disorders like platelet aggregation; cancer, including, *inter alia*, colorectal, lung and prostate cancers; digestive

15 disorders, such as colitis, Crohn's disease and inflammatory bowel disease; eye conditions, e.g., uveitis, cataracts; heart disease, e.g., acute myocardial, cardiovascular thrombosis; infectious diseases, e.g., sepsis, osteomyelitis; neurologic

20 disorders, e.g., multiple sclerosis, stroke; respiratory diseases, e.g., allergic asthma, allergic rhinitis; skin disorders, e.g., psoriasis; transplantation problems, e.g., graft-versus-host disease, organ transplant rejection; reduction in

25 sensitivity allergens, e.g., peanuts, and injuries resulting trauma etc.

In another embodiment, the crystallized whole antibodies, scFv antibody fragments and Fab antibody fragments, and compositions and formulations thereof,

30 according to this invention, may be used alone or in test kits to diagnose diseases or infections including, *inter alia*, osteomyelitis, salmonellosis, shigellosis, and the location and extent of disease staging in cancers such as non-Hodgkin's lymphoma and leukemia.



In yet another embodiment, the crystallized whole antibodies, scFv antibody fragments and Fab antibody fragments, and compositions and formulations thereof, according to this invention, may be used as *in vivo* imaging agents for the detection of diseases such as cardiovascular thrombosis.

The antibodies that may be crystallized and used according to this invention include, but are not limited to: anti-cytokine antibodies, anti-CD antigen antibodies (anti-CD3, -CD20 (e.g., Rituximab), anti-CD25, anti-CD52, anti-CD33, anti-CD11a), anti-TNF- $\alpha$  (e.g., Infliximab), anti-rattlesnake venom, anti-ICAM (e.g., anti-ICAM-1, anti-ICAM-3), anti-growth factor antibodies (e.g., anti-VEGF), anti-growth factor receptor antibodies (e.g., anti-HER2/neu (e.g., Trastuzumab), anti-EGFR), anti-immunoglobulin antibodies (e.g., anti-IgE), anti-polyclonal Ab antibodies, anti-viral antibodies (e.g., anti-CMV, anti-HIV (e.g., anti-gp120), anti-HBV, anti-RSV (e.g., anti-F glycoprotein)), etc.), anti-complement antibodies (e.g., anti-C5), anti-clotting factor antibodies (e.g., anti-gpIIb/IIIa, anti-Factor VII), anti-interleukin antibodies (e.g., anti-IL-5, anti-IL-4, anti-IL-8), antibodies targeted to the Major Histocompatibility Complex (e.g., anti-HLA), anti-idiotypic antibodies, anti-integrin antibodies (e.g., anti- $\beta$ -2-integrin), anti-17-IA cell surface antigen, anti- $\alpha$ 4 $\beta$ 7, anti-VLA-4, and anti-CBL.

Those of skill in the art will appreciate that antibody fragments including, *inter alia*, Fv and Fab antibody fragments of the above-mentioned whole antibodies may also be crystallized and used according to this invention.

Production of crystals of a whole antibody or crystals of a single-chain Fv antibody fragment, or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals

5                   According to the one embodiment of this invention, crystals of a whole antibody, crystals of a single-chain Fv antibody fragment, crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals are prepared by the following  
10 process.

                  First, the whole antibody or single-chain Fv antibody fragment or Fab antibody fragment is crystallized. Next, excipients or ingredients selected from sugars, sugar alcohols, viscosity increasing  
15 agents, wetting or solubilizing agents, buffer salts, emulsifying agents, antimicrobial agents, antioxidants, and coating agents are added directly to the mother liquor. Alternatively, the mother liquor is removed, after which the crystals are suspended in an excipient  
20 solution for a minimum of 1 hour to a maximum of 24 hours. The excipient concentration is typically between about 0.01 and about 10% (w/w). The ingredient concentration is between about 0.01 and about 90% (w/w). The crystal concentration is between about 0.01  
25 and about 99% (w/w).

                  The mother liquor is then removed from the crystal slurry either by filtration or by centrifugation. Subsequently, the crystals are washed optionally with solutions of about 50 to 100% (w/w) of  
30 one or more organic solvents such as, for example, ethanol, methanol, isopropanol or ethyl acetate, either at room temperature or at temperatures between -20 °C to 25 °C.

The crystals are then dried either by passing a stream

of nitrogen, air, or inert gas over them. Alternatively, the crystals are dried by air drying, spray drying, lyophilization or vacuum drying. The drying is carried out for a minimum of about 1 hour to  
5 a maximum of about 72 hours after washing, until the moisture content of the final product is below about 10% by weight, most preferably below about 5% by weight. Finally, micromizing (reducing the size) of the crystals can be performed if necessary.

10               According to one embodiment of this invention, when preparing crystals of a whole antibody, or crystals of a single-chain Fv antibody fragment, or crystals of an Fab antibody fragment, or formulations or compositions comprising such crystals, enhancers,  
15 such as surfactants, are not added during crystallization. Excipients or ingredients are added to the mother liquor after crystallization, at a concentration of between about 1 and about 10% (w/w), alternatively at a concentration of between about 0.1  
20 and about 25% (w/w), alternatively at a concentration of between about 0.1 and about 50% (w/w). The excipient or ingredient is incubated with the crystals in the mother liquor for about 0.1 to about 3 hrs, alternatively the incubation is carried out for about  
25 0.1 to about 12 hrs, alternatively the incubation is carried out for about 0.1 to about 24 hrs.

              In another embodiment of this invention, the ingredient or excipient is dissolved in a solution other than the mother liquor, and the crystals are  
30 removed from the mother liquor and suspended in the excipient or ingredient solution. The ingredient or excipient concentrations and the incubation times are the same as those described above.

              The present invention may also utilize other

slow release methodologies, such as silicon based rings or rods which have been preloaded with encapsulated crystals of a whole antibody or crystals of a single-chain Fv antibody fragment, or crystals of an Fab  
5 antibody fragment, or formulations or compositions comprising them, and can therefore act as implants for delivery. Such methodologies provide a constant level of antibodies or antibody fragments to the bloodstream over a period of weeks or months. Such implants can be  
10 inserted intradermally and can be safely replaced and removed when needed.

Other formulations and compositions according to this invention include vaccine formulations and compositions comprising crystals of a whole antibody,  
15 or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, and adjuvant and/or encapsulating polymer(s). In one embodiment of this invention, a whole anti-idiotypic antibody is itself the immunogen. In this embodiment, the whole  
20 antibody crystals and crystal formulations or compositions would elicit a response to the antigen that the anti-idiotypic mimics or is closely related to. Therefore, the anti-idiotypic antibody can act as a type of vaccine or therapy against cancers and  
25 autoimmune diseases, e.g., allergies, as well as viruses, for example, hepatitis B virus.

One embodiment of such vaccine formulations or compositions involves a single vaccine injection containing microspheres comprising crystalline whole  
30 antibodies, or scFv fragments or Fab fragments thereof. Those microspheres would, for example, be characterized by three or more different release profiles. In this way, crystals of a whole antibody, or fragment thereof, that act like antigens may be released over a sustained

period sufficient to generate lasting immunity. By virtue of such a formulation or composition, multiple antigen boosts may be made available in single unit form. One advantage of such a system is that by using  
5 whole antibody crystals, crystals of single-chain Fv antibody fragments, or crystals of Fab antibody fragments, or formulations or compositions comprising such crystals, the native three-dimensional structures of the antibodies or antibody fragments are maintained  
10 and presented to the immune system in their native form, thus eliciting the immune response seen with native antibodies.

Once the immune system is primed, there may be less need for an adjuvant effect. Therefore, in the  
15 slower degrading inoculations, a less immunogenic adjuvant may be included and possibly no adjuvant may be required in the slowest degrading microspheres of the formulations and compositions. In this way, patient populations in remote areas would not have to  
20 be treated multiple times in order to provide protection against infectious diseases.

Another advantage of the present invention is that crystals of a whole antibody or crystals of a single-chain Fv antibody fragment, or crystals of an  
25 Fab antibody fragment, or formulations thereof, that are encapsulated within polymeric carriers to form compositions comprising microspheres can be dried by lyophilization. Lyophilization, or freeze-drying allows water to be separated from the composition. The  
30 antibody crystal composition is first frozen and then placed in a high vacuum. In a vacuum, the crystalline H<sub>2</sub>O sublimates, leaving the whole antibody crystal or antibody fragment crystal composition behind, containing only the tightly bound water. Such

processing further stabilizes the composition and allows for easier storage and transportation at typically encountered ambient temperatures.

Spray drying allows water to be separated  
5 from the crystal preparation. It is highly suited for the continuous production of dry solids in either powder, granulate or agglomerate form from liquid feedstocks as solutions, emulsions, and pumpable suspensions. Spray drying involves the atomization of  
10 a liquid feedstock into a spray of droplets and contacting the droplets with hot air in a drying chamber. The sprays are produced by either rotary (wheel) or nozzle atomizers. Evaporation of moisture from the droplets and formation of dry particles  
15 proceed under controlled temperature and airflow conditions. Relatively high temperatures are needed for spray drying operations. However, heat damage to products is generally only slight, because of an evaporative cooling effect during the critical drying  
20 period and because the subsequent time of exposure to high temperatures of the dry material may be very short. Powder is discharged continuously from the drying chamber. Operating conditions and dryer design are selected according to the drying characteristics of  
25 the product and the powder specification. Spray drying is an ideal process where the end product must comply with precise quality standards regarding particle size distribution, residual moisture content, bulk density and particle shape.

30 This feature is especially desirable for therapeutic antibodies and anti-idiotypic vaccines, which can be dispensed into single dose sterile containers ("ampules") or, alternatively, any desired increment of a single dose as a slurry, in a

formulation or a composition. The ampules containing the dispensed slurries, formulations or compositions can then be capped, batch frozen and lyophilized under sterile conditions. Such sterile containers can be  
5 transported throughout the world and stored at ambient temperatures. Such a system is useful for providing sterile vaccines and therapeutic antibodies to remote and undeveloped parts of the world. At the point of use, the ampule is rehydrated with the sterile solvent  
10 or buffer of choice and dispensed. For such preparations, minimal or no refrigeration is required.

In another embodiment of this invention, crystals of a whole antibody, or crystals of a single-chain Fv antibody fragment, or crystals of an Fab  
15 antibody fragment, according to this invention may be crosslinked for additional stability. This is advantageous for the use of such crystals, crystal formulations and compositions in areas of pH extremes, such as the gastrointestinal tract of humans and  
20 animals. For example, antibody crystals, such as, monoclonal antibody crystals, may be crosslinked using one of a variety of crosslinkers, including, but not limited to, Dimethyl 3, 3'-dithiobispropionimide.HCl (DTBP), Dithiobis (succinimidylpropionate) (DSP), Bis  
25 maleimido- hexane (BMH), Bis[Sulfosuccinimidyl]suberate (BS), 1,5-Difluoro-2,4-dinitrobenzene (DFDNB), Dimethylsuberimide.2HCl (DMS), Disuccinimidyl glutarate (DSG), Disulfosuccinimidyl tartarate (Sulfo-DST), 1-Ethyl-3-  
30 [3-Dimethylaminopropyl]carbodiimide hydrochloride (EDC), Ethylene glycolbis[sulfosuccinimidylsuccinate] (Sulfo-EGS), N-[g-maleimidobutyryloxy]succinimide ester (GMBS), N-hydroxysulfosuccinimidyl-4-azidobenzoate (Sulfo-HSAB), Sulfosuccinimidyl-6-[a-methyl-a-(2-

pyridyldithio)toluamido] hexanoate (Sulfo-LC-SMPT),  
Bis-[b-(4-azidosalicylamido) ethyl]disulfide (BASED)  
and glutaraldehyde (GA).

In a further embodiment of this invention,  
5 crystals of a whole antibody or scFv fragment of an  
antibody or Fab fragment of an antibody may be  
radiolabelled to be used in antibody radiation  
therapies. In such a therapy, for example, a  
radiolabelled anti-cancer antibody crystal or scFv  
10 fragment crystal or Fab antibody fragment crystal, or  
formulation or composition comprising such crystals,  
can be delivered according to this invention, to the  
site of the cancer. After delivery, the released  
antibody or scFv fragment or Fab antibody fragment  
15 binds to its targeted cancer antigen and delivers the  
radioisotope directly to the cancerous cells or tumor.  
The release of the antibody may be timed according to  
this invention. Alternatively, when using crosslinked  
crystals in radiation therapy, the crosslinkers  
20 themselves may be radiolabeled. In this embodiment,  
the whole antibody, Fv antibody fragment or Fab  
antibody fragment that is in the crosslinked crystal  
serves to target and deliver the radioisotope to the  
cancerous cell or tumor. The radioisotope itself is  
25 carried and released by the crosslinker.  
Theoretically, useful radiolabels include, but are not  
limited to, the following radioisotopes or  
radionucleotides:  $^3\text{H}$ ,  $^{14}\text{C}$ ,  $^{15}\text{N}$ ,  $^{35}\text{S}$ ,  $^{90}\text{Y}$ ,  $^{99}\text{Tc}$ ,  $^{111}\text{In}$ ,  $^{125}\text{I}$ ,  
 $^{131}\text{I}$ . Practically, however, *in vivo* use in  
30 radiotherapies would limit the radiolabel to  $^{131}\text{I}$ ,  $^{90}\text{Y}$ ,  
or any other radiolabels defined by a short half-life.  
For example, the monoclonal antibody Rituximab (see  
Example 1) has been labeled with  $^{90}\text{Yttrium}$  ( $^{90}\text{Y}$ ), in  
order to be used for radioimmunotherapy in patients



with non-Hodgkin's lymphomas. This compound is commercially available as Ibritumomab tiuxetan (Zevalin<sup>™</sup>) (IDEC Pharmaceuticals, (San Diego, CA).

Batch Crystallization of Crystals of a Whole Antibody  
5 or Crystals of a Single-Chain Fv Antibody Fragment or  
Crystals of an Fab Antibody Fragment

Protein crystals are grown by controlled crystallization of protein from aqueous solutions or aqueous solutions containing organic solvents.

10 Solution conditions that may be controlled include, for example, the rate of evaporation of solvent, organic solvents, the presence of appropriate co-solutes and buffers, pH and temperature. A comprehensive review of the various factors affecting the crystallization of

15 proteins has been published by McPherson, A., Methods Enzymol. 114:112-20 (1985).

Large-batch (industrial-scale) crystallization typically involves a much greater range of conditions than does crystallization by the

20 classical "hanging drop" method. The initial protein concentration ranges between about 1 and about 200 mg/ml (or possibly even more), more preferably from about 0.01 mg/ml to about 500 mg/ml, for large batch crystallization, while the protein concentration for

25 the hanging drop method is limited to about 4 to about 10 mg/ml (in rare cases, up to about 25 mg/ml). A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a

30 protein concentration that is within a range from about 0.01 mg/ml up to and including about 3.9 mg/ml. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment

thereof, wherein said crystal is crystallized using a protein concentration that is within a range from about 4 mg/ml up to and including about 10 mg/ml. A further embodiment of this invention is a crystal of a whole  
5 antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a protein concentration that is within a range from about 10.1 mg/ml up to and including about 25 mg/ml. A further embodiment of this invention is a crystal of a whole  
10 antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a protein concentration that is within a range from about 25.1 mg/ml up to and including about 200 mg/ml. A further embodiment of this invention is a crystal of a whole  
15 antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a protein concentration that is within a range from about 200.1 mg/ml up to about 500 mg/ml.

The crystallization buffers used for large-  
20 batch crystallization can have a pH range of about 3 to about 10, more preferably a pH range from about pH 1.9 to about pH 11.1, while the hanging drop method is carried out at a pH range of about 5.0 to about 9.0 (though usually this is accomplished at a pH at or  
25 around about 7.0). A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range that is from greater than about pH 1.9 up to and including about pH 2.9. A  
30 further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 2.9 up to and including about pH 3.9. A further embodiment of this

invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 3.9 up to and including about pH 4.9. A

5 further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 4.9 up to and including about pH 5.9. A further embodiment of this

10 invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 5.9 up to and including about pH 7.9. A further embodiment of this invention is a crystal of a

15 whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 7.9 up to and including about pH 8.9. A further embodiment of this invention is a crystal of a whole antibody, or an Fab

20 or scFv antibody fragment thereof, wherein said crystal is crystallized within a pH range from greater than about pH 8.9 up to and including about pH 9.9. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment

25 thereof, wherein said crystal is crystallized within a pH range from greater than about pH 9.9 up to and including about pH 11.1.

Large-batch crystallization can be accomplished at temperatures that range from about 4°C

30 to about 37°C, more preferably temperatures that range from about -21°C up to about +61°C, while most hanging drop crystallization is carried out at temperatures from about 4°C up to room temperature (about 22°C). A further embodiment of this invention is a crystal of a

whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a temperature range from about -21°C up to below about 4°C. A further embodiment of this invention is a

5 crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a temperature range from about 4°C up to and including room temperature (about 22°C). A further embodiment of this invention is a crystal of a whole

10 antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a temperature range from above room temperature (about 22°C) up to and including about 37°C. A further embodiment of this invention is a crystal of a whole

15 antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized within a temperature range from above about 37°C up to about 61°C.

A range of about 5% to about 40% polyethylene glycol (PEG), more preferably a PEG concentration from

20 glycol (PEG), more preferably a PEG concentration from about 2% to about 80%, with a PEG size (chain length, i.e., number of ethylene glycol residues in the PEG chain) of about 200 to about 20,000, more preferably a PEG size (chain length) of about 200 to about 40,000,

25 more preferably a PEG size (chain length) of about 200 to about 80,000, can be used in the crystallization buffers for large-batch crystallization.

Theoretically, hanging drop methods can also use these sizes and concentrations of PEG, but normally

30 conditions will not go outside a range of about 5% to about 20% PEG 400 (size (chain length)) to PEG 10000 (size (chain length)). A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal

is crystallized using a PEG concentration that is within a range from about 2% up to about 4.9%. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a PEG concentration that is within a range from about 5% up to and including about 20%. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a PEG concentration that is within a range from about 20.1% up to and including about 80%. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a PEG size (chain length) that is within a range from about 200 up to about 400. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a PEG size (chain length) that is within a range from about 400 up to and including about 10,000. A further embodiment of this invention is a crystal of a whole antibody, or an Fab or scFv antibody fragment thereof, wherein said crystal is crystallized using a PEG size (chain length) that is within a range from greater than about 10,000 up to and including about 80,000.

Large-batch crystallization can be accomplished at buffer concentrations that range from 0 mM (no buffer) to about 4 M, while most hanging drop crystallization is carried out at buffer concentrations from about 2 mM to about 1 M.

Large-batch crystallization can be accomplished at metal or non-metal ion concentrations that range from about 0 mM (no buffer) to about 4 M.

Examples of metal and non-metal ions include, *inter alia*, calcium, magnesium, manganese, copper, zinc, lithium, ammonium, iron, cobalt, cesium, cadmium, nickel, sodium and potassium.

5                Large-batch crystallization can be accomplished at salt concentrations that range from about 0 mM (no buffer) to about 4 M. Examples of suitable salts include, *inter alia*, chloride, acetate, sulfate, phosphate, nitrate, citrate, Tris, HEPES,  
10 cacodylate, imidazole, CHES, CAPS, MES, MOPS, tartarate, borate, carbonate/bicarbonate, fluoride, iodide, thiocyanate, formate, malonate, succinic acid, bicine and EDTA.

              It will be understood by those of skill in  
15 the art that the crystallization methods according to this invention can be accomplished using a wide variety of reagents, under a wide variety of crystallization conditions, including but not limited to:  
crystallization in the presence of various divalent or  
20 monovalent ions; crystallization in the presence of concentrations of divalent metal ions ranging from about 5 mM to about 500 mM; crystallization using various buffer salts including, but not limited to acetate, borate, carbonate, succinate, imidazole, Tris,  
25 HEPES, MOPs, Phosphate, CHES, and other biological buffers mentioned in the Sigma catalogue;  
crystallization using reagents including, *inter alia*, PEG monomethyl ether, MPD, ethoxyethanol, propanediol, organic solvents, sodium or potassium salts like  
30 sulfite (including other sodium salts), ammonium salts, lithium salts, PEG derivatives, or any other organic compounds; crystallization methods that are stationary or involve tumbling or mixing; crystallization in the presence or absence of

detergents or chelators.

Crystals are obtained in large-batch crystallization in about 3 hours to about 72 hours maximum, more preferably in about 5 minutes to about 48  
5 hours, while hanging drop methods can take days, weeks, or even months to yield crystals. Furthermore, large-batch crystallization uses an agitation step, unlike hanging drop protocols.

Large-batch crystallization is performed as  
10 follows: a suitable volume of the antibody or scFv fragment or Fab antibody fragment to be crystallized (in its storage buffer) is mixed with an equal volume of a crystallization solution or crystallization buffer (at a prescribed pH). The mixture can either be seeded  
15 with crushed crystals that were previously obtained (through other experiments) or used without seeding. The mixture is then tumbled, for example, in a hematology/chemistry mixer for about 3 to about 48 hours at the desired temperature (typically about room  
20 temperature).

Large-batch crystallization may or may not involve the use of "seed" crystals, i.e., crystals obtained during small-scale crystallization screens for the determination of crystallization conditions.  
25 Typically, seed crystals can be obtained from hanging-drop methods using commercially-available crystallization screening kits (e.g., Wizard I and Wizard II, and Cryo I and Cryo II kits (Emerald BioStructures, Inc. (Bainbridge Island, WA)), or  
30 Crystal Screen and Crystal Screen II kits (Hampton Research (Laguna Niguel, CA). Alternatively, crystals of a whole antibody or scFv fragment or Fab antibody fragment may be prepared using a screening method, called microbatch screening, which is, in practice, a

scaled down version of the large-batch crystallization method described above.

Although crystallization of whole antibodies has been a subject of significant interest for approximately the last twenty-five years, very few have been crystallized [Harris L.J., Skaletsky, E., and McPherson, A., J. Mol. Biol. 275:861-72 (1998); Harris L.J., Larson, S.B., Skaletsky, E., and McPherson, A., Immunological Reviews 163:35-43 (1998)]. Such prior efforts utilized solely hanging drop or seeding drop protocols. Both methods were characterized by extremely low yields of crystals and, therefore, were unsuitable for large-scale production of antibody crystals. Such was the case because of difficulties in antibody crystallization due to their large size, the presence of surface oligosaccharides, and their high degree of segmental flexibility. Crystallization of whole antibodies on a large scale, a process offering an alternative route of delivery for the therapeutic antibodies, has never been explored before.

Encapsulation of Crystals of A Whole Antibody or a Single-chain Fv Antibody Fragment or a Fab Antibody Fragment in Polymeric Carriers

According to one embodiment of this invention, compositions are produced when whole antibody crystals or crystals of a single-chain Fv antibody fragment, or crystals of an Fab antibody fragment, or formulations comprising such crystals, are encapsulated in at least one polymeric carrier to form microspheres by virtue of encapsulation within the matrix of the polymeric carrier to preserve their native and biologically active tertiary structure. The crystals can be encapsulated using various



biocompatible and/or biodegradable polymers having unique properties which are suitable for delivery to different biological environments or for effecting specific functions. The rate of dissolution and, therefore, delivery of active antibodies or fragments thereof is determined by the particular encapsulation technique, polymer preparation, polymer crosslinking, polymer thickness, polymer solubility, and antibody crystal geometry.

Crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment, or formulations of such crystals, to be encapsulated are suspended in a polymeric carrier which is dissolved in an organic solvent. The polymer solution must be concentrated enough to completely coat the antibody crystals or formulations after they are added to the solution. Such an amount is one which provides a weight ratio of antibody crystals to polymer between about 0.02 and about 20, preferably between about 0.1 and about 2. The antibody crystals are contacted with polymer in solution for a period of time between about 0.5 minutes and about 30 minutes, preferably between about 1 minute and about 3 minutes. The crystals should be kept suspended and not allowed to aggregate as they are coated by contact with the polymer.

Following that contact, the crystals become coated and are referred to as nascent microspheres. The nascent microspheres increase in size during the coating process. In a preferred embodiment of the invention, the suspended coated crystals or nascent microspheres along with the polymeric carrier and organic solvent are transferred to a larger volume of an aqueous solution containing a surface active agent,

known as an emulsifier. In the aqueous solution, the suspended nascent microspheres are immersed in the aqueous phase, where the organic solvent evaporates or diffuses away from the polymer. Eventually, a point is reached where the polymer is no longer soluble and forms a precipitated phase encapsulating the antibody crystals, antibody fragment crystals, or formulations to form a composition. This aspect of the process is referred to as hardening of the polymeric carrier or polymer. The emulsifier helps to reduce the interfacial surface tension between the various phases of matter in the system during the hardening phase of the process. Alternatively, if the coating polymer has some inherent surface activity, there may be no need for addition of a separate surface active agent.

Emulsifiers useful to prepare encapsulated crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment according to this invention include poly(vinyl alcohol) as exemplified herein, surfactants and other surface active agents which can reduce the surface tension between the polymer coated whole antibody crystals or polymer coated crystal formulations and the solution.

In a preferred embodiment of this invention, crystallinity of the whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment is maintained during the encapsulation process. The crystallinity is maintained during the coating process by using an organic solvent in which the crystals are not soluble. Subsequently, once the coated crystals are transferred to the aqueous solvent, rapid hardening of the polymeric carrier and sufficient coating of the

crystals in the previous step shields the crystalline material from dissolution.

The polymers used as polymeric carriers to coat the whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment can be either homo-polymers or co-polymers. The rate of hydrolysis of the microspheres is largely determined by the hydrolysis rate of the individual polymer species. In general, the rate of hydrolysis decreases as follows: polycarbonates > polyesters > polyurethanes > polyorthoesters > polyamides. For a review of biodegradable and biocompatible polymers, see W.R. Gombotz and D.K. Pettit, "Biodegradable polymers for protein and peptide drug delivery", Bioconjugate Chemistry, vol. 6, pp. 332-351 (1995).

In a preferred embodiment of this invention, the polymeric carrier comprises a single polymer type, such as PLGA. In a next preferred embodiment, the polymeric carrier can be a mixture of polymers, such as 50% PLGA and 50% albumin.

Other polymers useful as polymeric carriers to prepare encapsulated crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment according to this invention include biocompatible/biodegradable polymers selected from the group consisting of poly (acrylic acid), poly (cyanoacrylates), poly (amino acids), poly (anhydrides), poly (depsipeptide), poly (esters), such as poly (lactic acid) or PLA, poly (b-hydroxybutyrate), poly (caprolactone) and poly (dioxanone); poly (ethylene glycol), poly (hydroxypropyl) methacrylamide, poly [(organo)phosphazene], poly (ortho esters), poly (vinyl alcohol), poly (vinylpyrrolidone), maleic

anhydride-alkyl vinyl ether copolymers, pluronic polyols, albumin, alginate, cellulose and cellulose derivatives, starch and its derivatives, collagen, fibrin, gelatin, hyaluronic acid, oligosaccharides, glycaminoglycans, sulfated polysaccharides, blends and copolymers thereof. Other useful polymers are described in J. Heller and R.W. Balar, "Theory and Practice of Controlled Drug Delivery from Biodegradable Polymers," Academic Press, New York, NY, (1980); K.O.R. Lehman and D.K. Dreher, Pharmaceutical Technology, vol. 3 (1979); E.M. Ramadan, A. El-Helw and Y. El-Said, Journal of Microencapsulation, vol. 4, pp. 125-132 (1987); O. Phillai and R. Panchagnula, Current Opinion in Chemical Biology, vol. 5, pp. 447-451 (2001). The preferred polymer will depend upon the particular antibody crystals or antibody fragment crystals used and the intended use of the encapsulated crystals or crystal formulations. Alternatively, the solvent evaporation technique may be used for encapsulating whole antibody crystals or antibody fragment crystals (see D. Babay, A. Hoffmann and S. Benita, Biomaterials vol. 9, pp. 482-488 (1988)).

In a preferred embodiment of this invention, crystals of a whole antibody or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment are encapsulated in at least one polymeric carrier using a double emulsion method, as illustrated herein, using a polymer, such as polylactic-co-glycolic acid. In a most preferred embodiment of this invention, the polymer is Polylactic-co-glycolic acid ("PLGA"). PLGA is a copolymer prepared by polycondensation reactions with lactic acid ("L") and glycolic acid ("G"). Various ratios of L and G can be used to modulate the

crystallinity and hydrophobicity of the PLGA polymer. Higher crystallinity of the polymer results in slower dissolution. PLGA polymers with 20-70% G content tend to be amorphous solids, while high level of either G or  
5 L result in good polymer crystallinity. See D.K. Gilding and A.M. Reed, "Biodegradable polymers for use in surgery-poly(glycolic)/poly(lactic acid) homo and copolymers: 1., *Polymer* vol. 20, pp. 1459-1464 (1981). PLGA degrades after exposure to water by hydrolysis of  
10 the ester bond linkage to yield non-toxic monomers of lactic acid and glycolic acid.

Another embodiment of this invention includes double-walled polymer coated microspheres. Double-walled polymer coated microspheres may be produced by  
15 preparing two separate polymer solutions in methylene chloride or other solvent which can dissolve the polymers. The whole antibody crystals or crystals of a single-chain Fv antibody fragment or crystals of an Fab antibody fragment are added to one of the solutions and  
20 dispersed. Here, the crystals become coated with the first polymer. Then, the solution containing the first polymer coated crystals is combined with the second polymer solution. [See Pekarek, K.J.; Jacob, J.S. and Mathiowitz, E. Double-walled polymer microspheres for  
25 controlled drug release, *Nature*, 367, 258-260]. As a result, the second polymer encapsulates the first polymer which is encapsulating the crystal. Ideally, this solution is then dripped into a larger volume of an aqueous solution containing a surface active agent  
30 or emulsifier. In the aqueous solution, the solvent evaporates from the two polymer solutions and the polymers are precipitated.

Whole antibodies and single-chain Fv antibody fragments and Fab antibody fragments recovered

by dissolving crystals, formulations or compositions according to this invention may be characterized for secondary structure. Such whole antibody crystals or crystals of a single-chain Fv antibody fragment or  
5 crystals of an Fab antibody fragment may also be characterized by  $\beta$ -sheet structural content, as indicated by a correlation spectra as compared to the spectra of the soluble antibody or antibody fragment counterpart determined by Fourier transform infrared  
10 (FTIR) spectra that is between about 0.8 and about 1.0. A correlation coefficient of less than about 0.8 indicates an protein sample that has become denatured to such an extent that its intermolecular  $\beta$ -sheet secondary structure content has increased, resulting in  
15 protein aggregation and precipitation.

In order that this invention may be better understood, the following examples are set forth. These examples are for the purpose of illustration only and are not to be construed as limiting the scope of  
20 the invention in any manner.

#### EXAMPLES

In the following examples, the crystal screening kits used (when seeding from hanging drops) were one or more of the following: Wizard I and Wizard  
25 II, and Cryo I and Cryo II kits (Emerald BioStructures, Inc. (Bainbridge Island, WA), or Crystal Screen and Crystal Screen II kits (Hampton Research (Laguna Niguel, CA). Batch crystallization or microbatch screening of whole antibodies was carried out by mixing  
30 the antibody with the appropriate crystallization buffer, followed by tumbling or incubation (with or without shaking as noted). Antibody crystals, which were obtained from vapor diffusion hanging drops in

preliminary screening or from microbatch screening, were used in most cases, to facilitate the crystallization process. Antibody crystals were confirmed by determining their birefringence.

5    Example 1            Rituximab

Rituximab is a chimeric murine/human monoclonal antibody commercially available as Rituxan™ (Genentech, Inc., South San Francisco, CA). This monoclonal antibody has been widely used to treat non-  
10    Hodgkins lymphoma. Rituximab is a chimeric IgG1 kappa immunoglobulin that binds to the CD20 antigen on the surface of normal and malignant B-lymphocytes. It is composed of murine light- and heavy-chain variable region sequences and a human constant region sequence.  
15    The Rituximab antibody has an approximate molecular weight (MW) of 145 kD.

Rituximab crystallization, batch 1:

Materials:

A - Rituximab antibody (stored until use  
20    at 4°C, at 10 mg/ml in 9.0 mg/ml sodium chloride, 7.35 mg/ml sodium citrate anhydrate, 0.7 mg/ml Polysorbate 80 and sterile water, pH 6.5)

B - Wizard I crystal screening kit  
(Emerald BioStructures, Bainbridge Island, WA)

25                    C - Polyethylene glycol-1000 (PEG-1000)

D - Imidazole

E - Calcium acetate buffer pH 8.0

Procedure:

Rituximab seed crystals were obtained from  
30    vapor diffusion drops in a preliminary screening using the Wizard I screening kit. Microbatch crystallization was carried out using 500 µl of a crystallization buffer containing 20% (w/v) PEG-1000, 100 mM imidazole,

and 200 mM calcium acetate, pH 8.0. After seeding with Rituximab seed crystals from the hanging drops, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific, Pittsburgh, PA) at 50 rpm at room temperature overnight. The antibody crystals obtained from this step were used as seeds for large-batch crystallization, which is, in essence, a scaled-up microbatch procedure. Large-batch crystallization was initiated by mixing 8 ml of the Rituximab solution with 8 ml of a crystallization buffer containing 20% (w/v) PEG-1000, 100 mM imidazole, and 200 mM calcium acetate, pH 8.0. The final concentration of the Rituximab in solution was 5 mg/ml. After seeding with Rituximab seed crystals from the microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After overnight tumbling, Rituximab crystals in the form of needle clusters were formed. 85% of the input Rituximab was crystallized in this example.

#### Example 2

##### Rituximab crystallization, batch 2:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 500  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization buffer containing 20% (w/v) PEG-1000, 100 mM imidazole, and 200 mM calcium acetate, pH 7.0. After seeding with Rituximab seed crystals from a microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After overnight tumbling, Rituximab crystals in the form of needle clusters were formed. This protocol has been



repeated, at pH 4.0, 5.0, 6.0 and 8.0, with similar results.

### Example 3

#### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 60  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization buffer containing 20% (w/v) PEG-600, 100 mM calcium acetate, and 50 mM 2-[N-cyclohexylamino] ethanesulfonic acid (CHES), pH 9.5. The final concentration of the Rituximab in solution was 5 mg/ml. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was incubated in a benchtop shaker/incubator (New Brunswick Scientific, Model C25), at 25°C at 650 rpm. After overnight incubation, Rituximab crystals in the form of needle clusters (Figure 1) were formed. 80% of the input Rituximab was crystallized in this example.

### Example 4

#### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 50  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization solution containing 20% (w/v) PEG-1000 and 100 mM calcium acetate. The mixture was allowed to sit at room temperature for 12 hours. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was allowed to continue incubating at room temperature in Eppendorf

centrifuge tubes. After overnight incubation, Rituximab crystals in the form of needle clusters were formed. This screen was repeated, with the calcium acetate concentration being adjusted to 10, 20, 40, 60, 80, 200 or 400 mM, with crystals being obtained under all conditions tested.

#### Example 5

##### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 50  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization solution containing 25% (w/v) PEG-1000 and 100 mM calcium acetate. The mixture was allowed to sit at room temperature for 12 hours. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was allowed to continue incubating at room temperature in centrifuge tubes. After overnight incubation, Rituximab crystals in the form of needle clusters were formed. This screen was repeated, with the PEG-1000 concentration being adjusted to 5, 10, 15, 20 or 40% (w/v), with crystals being obtained under all conditions being tested.

#### Example 6

##### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 50  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization buffer containing 20% (w/v) PEG-6000, 100 mM calcium acetate and 100 mM Tris, pH

8.0. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was incubated in a benchtop shaker/incubator (New Brunswick Scientific, Model C25), at 25°C at 225 rpm. After  
5 overnight incubation, Rituximab crystals in the form of needle clusters were formed. This example was repeated, by substituting PEG-2000, PEG-4000 or PEG-8000 for the PEG-6000 (maintaining the PEG concentration at 20%), with crystals being obtained  
10 under all conditions being tested. See Figure 1.

#### Example 7

##### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in  
15 Example 1, except that crystallization was initiated by mixing 60  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization buffer containing 20% (w/v) PEG-6000, 100 mM calcium acetate and 100 mM Tris, pH 7.0. After seeding with Rituximab seed crystals from a  
20 previously obtained microbatch, the mixture was incubated in a benchtop shaker/incubator (New Brunswick Scientific, Model C25), at 25°C at 650 rpm. After overnight incubation, Rituximab crystals in the form of needle clusters were formed. This screen was repeated,  
25 by substituting PEG-200, PEG-300 or PEG-20000 for PEG-6000 (maintaining the PEG concentration at 20%). 140 mM (instead of 100 mM) Tris was used for the screens using PEG-200. Crystals being obtained under all conditions tested.

#### 30 Example 8

##### Microbatch crystallization screening of Rituximab:

Rituximab crystals were obtained as in Example 1, except that crystallization was initiated by mixing 50  $\mu$ l of Rituximab (10 mg/ml) with an equal volume of crystallization buffer containing 20% (w/v) PEG-1000 and 200 mM  $\text{CuSO}_4$ , and 100 mM imidazole, pH 8.0. After sitting at room temperature for 14 hours, the mixture was seeded with Rituximab seed crystals from a previously obtained microbatch and incubated at room temperature. After overnight incubation, Rituximab crystals in the form of rods were formed. Subsequently, this screen was repeated, by substituting other divalent cations, e.g., 200 mM  $\text{CaCl}_2$ ,  $\text{MnCl}_2$ , or  $\text{ZnCl}_2$  for the 200 mM  $\text{CuSO}_4$ , with crystals in the form of needle clusters, disks and quasi crystals being obtained.

#### Example 9

##### Rituximab crystallization using dialyzed Rituximab

An aliquot of 4 ml of Rituximab (10 mg/ml) was dialyzed against 2 liters of deionized water overnight at 4°C, with two changes of deionized water before being concentrated to 1 ml with a centrifugal filter device (Millipore, 30 kD cut-off). Crystallization was carried out in hanging drops by mixing 6  $\mu$ l of concentrated Rituximab with 2  $\mu$ l of crystallization buffer containing 20% (w/v) PEG-1000 and 100 mM imidazole, 200 mM calcium acetate, pH 8.0, on a glass cover slide. The cover slide was flipped over and placed onto a well, which contained 450  $\mu$ l of the same buffer, in a 24-well plate. After incubation of the plate at room temperature for approximately one week, Rituximab crystals in the form of cubes were formed.

Example 10Rituximab crystallization using dialyzed  
Rituximab

An aliquot of 5 ml of Rituximab (10 mg/ml)  
5 was dialyzed against 2 liters of 10 mM Hepes buffer, pH  
7.0, overnight at 4°C, before being concentrated to 54  
mg/ml with a centrifugal filter device (Millipore, 10  
kD cut-off). Batch crystallization was carried out by  
mixing 20 µl of concentrated Rituximab with buffer and  
10 additives, so that the final mixture contained 36 mg/ml  
antibody, 133 mM Hepes, pH 7.50, 66 mM CaCl<sub>2</sub> and 13% 2-  
methyl-2,4,-pentanediol. The mixture was tumbled in a  
hematology/chemistry mixer (Model 346, Fisher  
Scientific) at room temperature at 50 rpm. After a 48  
15 hour incubation at room temperature, Rituximab crystals  
in the form of needle clusters were formed. In this  
example, 84% of the input Rituximab was crystallized.

Example 11Rituximab crystallization using dialyzed  
Rituximab

20 An aliquot of 5 ml of Rituximab (10 mg/ml)  
was dialyzed against 2 liters of 10 mM Hepes buffer, pH  
7.0, overnight at 4°C, before being concentrated to 54  
mg/ml with a centrifugal filter device (Millipore, 10kD  
25 cut-off). Batch crystallization was carried out by  
mixing 20 µl of concentrated Rituximab with buffer and  
additives, so that the final mixture contained 18 mg/ml  
antibody, 200 mM Hepes, pH 7.50, 200 mM CaCl<sub>2</sub> and 33%  
PEG-400. The mixture was tumbled in a  
30 hematology/chemistry mixer (Model 346, Fisher  
Scientific) at room temperature at 50 rpm. After a 48  
hour incubation, Rituximab crystals in the form of  
needle clusters were formed.

Example 12Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 80  $\mu$ l of antibody with 20  $\mu$ l of 0.02 M  $\text{CaCl}_2$ , 0.1 M sodium acetate, pH 4.6, 30% 2-methyl-2,4-pentanediol and tumbling the mixture in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After a 48 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

Example 13Rituximab crystallization in the presence of a detergent:

Rituximab was crystallized by mixing 450  $\mu$ l of Rituximab with an equal volume of a crystallization buffer containing 20% (w/v) PEG-1000, 100 mM imidazole, 200 mM calcium acetate, pH 8.0, and 0.1% Tween<sup>®</sup>80 (a detergent) (Sigma-Aldrich). After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at 50 rpm at room temperature. Crystals, in the form of needle clusters, were formed after overnight tumbling. Rituximab was also crystallized using starting volumes ranging between 50  $\mu$ l and 1.0 ml. The same type of crystals formed.

Example 14Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 50  $\mu$ l of antibody with 50  $\mu$ l of 0.2 M calcium acetate, 0.1 M sodium acetate, pH 4.6, 30% PEG 400.

After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24  
5 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

#### Example 15

##### Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by  
10 mixing 50 µl of antibody with 50 µl of 0.2 M calcium acetate, 0.1 M imidazole, pH 8.0, 10% PEG 8000. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a  
15 hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

#### Example 16

##### Rituximab crystallization:

20 Rituximab (10 mg/ml) was crystallized by mixing 50 µl of antibody with 50 µl of 0.2 M calcium acetate, 0.1 M Tris, pH 7.0, 20% PEG 3000. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a  
25 hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

#### Example 17

30 Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 50 µl of antibody with 50 µl of 0.2 M calcium acetate, 0.1 M MES, pH 6.0, 20% PEG 8000. After seeding with Rituximab seed crystals from a previously  
5 obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

10 Example 18

Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 50 µl of antibody with 50 µl of 0.05 M calcium acetate, 0.1 M imidazole, pH 8.0, 35% 2-ethoxyethanol.  
15 After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals  
20 in the form of needle clusters were formed.

Example 19

Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 50 µl of antibody with 50 µl of 0.05 M calcium  
25 acetate, 0.1 M acetate, pH 4.5, 40% 1,2-propanediol. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24  
30 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.



Example 20Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 50 µl of antibody with 50 µl of 0.2 M calcium acetate, 0.1 M HEPES, pH 7.5, 40% PEG 400. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

Example 21Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 4 ml antibody with 4 ml reagent containing 0.2 M calcium acetate, 0.1 M TRIS, pH 7.0, 20% PEG 6000 and 0.1% Tween®80 (Sigma-Aldrich). After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed. The yield was 93% by this method.

Example 22Rituximab crystallization:

Rituximab (10 mg/ml) was crystallized by mixing 1 ml antibody with 1 ml reagent containing 0.2 M calcium acetate, 0.1 M MES, pH 6.0, 20% PEG 6000 and 0.1% Tween®80 (Sigma-Aldrich). After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a

hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed. The yield was 80% by this method.

#### Example 23

##### Rituximab crystallization:

Dialyzed Rituximab (10 mg/ml) was crystallized by mixing 10  $\mu$ l of antibody with 10  $\mu$ l of 0.2 M calcium acetate, 0.1 M sodium acetate, pH 4.6, 30% 2-propanol. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

#### Example 24

##### Rituximab crystallization:

Dialyzed Rituximab (10 mg/ml) was crystallized by mixing 10  $\mu$ l of antibody with 20  $\mu$ l of 0.02 M  $\text{CaCl}_2$ , 0.1 M sodium acetate, pH 4.6, 30% 2-methyl-2,4-pentanediol. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

#### Example 25

##### Rituximab crystallization:

Dialyzed Rituximab (10 mg/ml) was crystallized by mixing 10  $\mu$ l of antibody with 20  $\mu$ l of 0.2 M  $\text{CaCl}_2$ , 0.1 M HEPES, pH 7.5, 28% PEG 400. After seeding with Rituximab seed crystals from a previously obtained microbatch, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of needle clusters were formed.

10 Example 26

Rituximab crystallization -- Different Crystal Form:

Rituximab (20 mg/ml) was crystallized by mixing 10  $\mu$ l of antibody with 10  $\mu$ l of solution containing 15% PEG 400, 0.51 M sodium sulfate, 0.1 M EDTA. The final concentration of the Rituximab in solution was 10.0 mg/ml. After seeding with Rituximab seed crystals from a previously obtained hanging drop, the mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of small needles were formed. In this example, 87% of the input Rituximab was crystallized. See Figure 4.

25 Example 27

Rituximab crystallization -- Different Crystal Form:

Rituximab (20 mg/ml) was crystallized by mixing 10  $\mu$ l of antibody with 10  $\mu$ l of solution containing 12% PEG 400 and 1.36 M sodium sulfate and 0.1 M Tris, pH 7.5. After seeding with Rituximab seed crystals from a previously obtained microbatch, the

mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 225 rpm. After a 24 hour incubation at room temperature, Rituximab crystals in the form of small  
5 needles were formed.

#### Example 28

##### Rituximab crystallization -- Different Crystal Form:

Dialyzed Rituximab (66 mg/ml) was  
10 crystallized by mixing 10 µl of antibody with 20 µl of solution containing of 0.2 M CaCl<sub>2</sub>, 0.1 M HEPES, pH 7.5, 28% PEG 400. After 20 days, additional amounts of PEG 400 (30 µl 100% PEG 400) and 10 µl of 1M lithium sulfate were added. After a 24 hour incubation at room  
15 temperature, Rituximab crystals in the form of cubes were formed. (Figure 3).

#### Example 29

##### Morphology of the crystals #1

Different forms of Rituximab were obtained by  
20 using different crystallization conditions.

For example, see Rituximab crystals from Figures 1 and 3.

#### Example 30

##### Morphology of the crystals #2

25 Crystallization:

Buffer: 100 mM Hepes, pH 7.7, 12% PEG 400, 1.17 M sodium sulfate.

Method: 1 volume Rituximab was mixed with 2 volumes crystallization buffer. The mixture was  
30 maintained at room temperature without agitation until crystals formed.

Result: Small (length  $\leq$  10  $\mu$ m) needle-like crystals formed. (Figure 4).

Example 31      Trastuzumab

Trastuzumab is a recombinant DNA-derived humanized monoclonal antibody commercially available as Herceptin™ (Genentech, Inc., South San Francisco, CA). This monoclonal antibody has been widely used to treat breast cancer which over-expresses the extracellular domain of the epidermal growth factor receptor 2 protein, HER2. Trastuzumab is an IgG1 kappa that contains human framework regions with the complementarity-determining regions of a murine antibody (4D5) that binds to HER2.

Trastuzumab Microbatch crystallization, batch 1:

Trastuzumab antibody was stored in its original 440 mg vial as a sterile lyophilized powder and was subsequently dissolved in 20 ml of sterile water. The dissolved Trastuzumab solution, containing 22 mg/ml Trastuzumab, 9.9 mg L-histidine HCl, 6.4 mg L-histidine, 400 mg  $\alpha,\alpha$ -trehalose dihydrate, and 1.8 mg polysorbate 20, USP.

A 210  $\mu$ l aliquot of Trastuzumab (22 mg/ml), in a buffer containing 0.495 mg/ml L-histidine HCl, 0.32 mg/ml L-histidine, 20 mg/ml  $\alpha,\alpha$ -trehalose dihydrate, and 0.09 mg/ml polysorbate 20, USP, was mixed with 210  $\mu$ l of crystallization buffer containing 25% PEG 400, 5% PEG 8000, 100 mM Tris, pH 8.5, 10% propylene glycol, and 0.1% Tween®80 (Sigma-Aldrich) and incubated at room temperature overnight. The final concentration of the Trastuzumab in solution was 11 mg/ml. This mixture was then seeded with Trastuzumab

crystals obtained from hanging drop and tumbled at 50 rpm in a Hematology/Chemistry mixer (Model 346, Fisher Scientific) after being supplemented with 20  $\mu$ l of propylene glycol. Trastuzumab crystals were obtained  
5 on the following day. (Figure 5). 85% of the input Trastuzumab was crystallized by this method.

#### Example 32

##### Trastuzumab Microbatch crystallization, batch 2:

10 50  $\mu$ l of Trastuzumab (22 mg/ml) in a buffering solution containing 0.495 mg/ml L-histidine HCl, 0.32 mg/ml L-histidine, 20 mg/ml  $\alpha,\alpha$ -trehalose dihydrate, and 0.09 mg/ml polysorbate 20, USP, was mixed with 50  $\mu$ l of crystallization buffer containing  
15 20% PEG 300, 10% glycerol, 0.1 M Tris, pH 7, 10% PEG 8000 and incubated at room temperature overnight after being seeded with Trastuzumab crystals obtained from microbatch. Trastuzumab crystals with a size ranging from 50-120  $\mu$ m were obtained on the following day.  
20 (Figure 6).

#### Example 33

##### Trastuzumab Microbatch crystallization, batch 3:

25 50  $\mu$ l of Trastuzumab (22 mg/ml), in a buffer containing 0.495 mg/ml L-histidine HCl, 0.32 mg/ml L-histidine, 20 mg/ml  $\alpha,\alpha$ -trehalose dihydrate, and 0.09 mg/ml polysorbate 20, USP, was mixed with 50  $\mu$ l of crystallization buffer containing 20% PEG 300, 10% glycerol, 0.1 M Tris, pH 7, 10% PEG 8000 and incubated  
30 at room temperature overnight after being seeded with Trastuzumab crystals obtained from microbatch.

Trastuzumab crystals with a size of about 20  $\mu\text{m}$  were obtained on the following day.

#### Example 34

##### Infliximab

5                   Infliximab is a chimeric murine/human monoclonal antibody commercially available as Remicade<sup>™</sup> (Centocor, Leiden, the Netherlands). This monoclonal antibody has been widely used to treat rheumatoid arthritis and Crohn's disease. Infliximab is a  
10 chimeric IgG1 kappa immunoglobulin that binds to the TNF- $\alpha$  antigen. It is composed of murine light- and heavy-chain variable region sequences and a human constant region sequence. The Infliximab antibody has an approximate molecular weight (MW) of 149 kD.

##### Infliximab Microbatch crystallization, batch 1:

                  Infliximab antibody was stored in its original 100 mg vial as a sterile lyophilized powder and was subsequently dissolved in 2 ml of sterile  
20 water. The dissolved solution, containing 100 mg Infliximab, 500 mg sucrose, 0.5 mg polysorbate 80, 2.2 mg monobasic sodium phosphate, and 6.1 mg dibasic sodium phosphate, was used for crystallization.

                  Infliximab was crystallized by mixing 50  $\mu\text{l}$   
25 of antibody (50 mg/ml) with 100  $\mu\text{l}$  of 35% ethoxyethanol, 0.2 M lithium sulfate, 0.1 M Tris, pH 8.6. The mixture was tumbled in a hematology/chemistry mixer (Model 346, Fisher Scientific) at room temperature at 50 rpm. After overnight incubation,  
30 Infliximab crystals in the form of rod shaped clusters were formed (Figure 2).

#### Example 35

Infliximab Microbatch crystallization,  
batch 2:

Rod shaped crystals similar to those of Example 34 were also obtained when Infliximab was  
5 incubated (without agitation) with 40% (w/v) PEG-400, 0.1 M Tris buffer, 200 mM lithium sulfate, pH 8.5, under the same conditions.

Example 36

Infliximab Microbatch crystallization,  
10 batch 3:

A 25 µl aliquot of Infliximab (50 mg/ml in 0.1M Tris HCl buffer, pH7.0) was mixed with 3 µl of 1M calcium chloride and 5 µl of 100% polyethylene glycol monomethyl ether 550 (PEG MME 550) and incubated  
15 (without agitation) overnight at room temperature. Cube shaped crystals of Infliximab formed overnight.

Example 37

Infliximab Microbatch crystallization

A 25 µl aliquot of Infliximab (20 mg/ml in  
20 water) was mixed with 50 µl of crystallization buffer containing 20% PEG 300, 0.1 M TRIS, pH 8.5 5% PEG 8000 and 10% glycerol. The final concentration of the Infliximab in solution was 6.7 mg/ml. This mixture was then incubated (without agitation) overnight at room  
25 temperature. Star shaped crystals of Infliximab formed after a week. (Figure 7).

Example 38

The crystallization conditions exemplified above are useful for the crystallization of any  
30 desirable clinically relevant antibody. Clinically relevant antibodies may be classified according to the



therapeutic area in which they are to be employed. Such antibodies include, but are not limited to, commercially available antibodies, including, but not limited to:

- 5 (1) Abciximab (ReoPro™) (anti-GPIIb/IIIa receptor; for the treatment of cardiovascular disease) (Centocor, Leiden, The Netherlands);
- (2) Palivizumab (Synagis™) (anti-F protein on RSV; respiratory disease) (manufactured by MedImmune
- 10 (Gaithersburg, MD);
- (3) Murumonab-CD3 (Orthoclone™) (anti-CD3 antibody; for tissue transplant rejection) (OrthoBiotech, Raritan, NJ);
- (4) Gemtuzumab ozogamicin (Mylotarg™) (cancer (anti-
- 15 CD33 antibody)) (Wyeth Labs, Philadelphia, PA);
- (5) Trastuzumab (Herceptin™) (cancer (anti-HER2 antibody)) (Genentech, South San Francisco, CA);
- (6) Basiliximab (Simulect™) (anti-CD25 antibody; for tissue transplant rejection) (Novartis, Basel,
- 20 Switzerland);
- (7) Daclizumab (Zenapax™) (anti-CD25 antibody; for tissue transplant rejection) (Protein Design Labs, Fremont, CA);
- (8) Etanercept (ENBREL™) (inflammatory disease)
- 25 (Immunex, Seattle, WA);
- (9) Ibritumomab tiuxetan (Zevalin™) (radioimmunotherapy for cancer) (IDEC Pharmaceuticals, San Diego, CA).

### Example 39

The crystallization conditions exemplified

30 above are useful for the crystallization of single-chain Fv (scFv) fragments of antibodies, or for crystallization of Fab fragments of antibodies.

Example 40Other Classes of Immunoglobulin

All embodiments of this invention are useful for crystallization of, and use of crystals of, all of the immunoglobulin classes IgG, IgM, IgA, IgD, IgE, and serum IgA (sIgA) as well as the subclasses IgG1, IgG2, IgG3 and IgG4, IgM1 and IgM2, and IgA1 and IgA2.

Example 41Crystallization as a Tool for MonoclonalAntibody Purification

Monoclonal antibodies including, *inter alia*, Rituximab, Infliximab and Trastuzumab may be obtained from mammalian cell culture. Crystallization of these monoclonal antibodies may be carried out either directly from the culture media or cell extract or after partial or complete purification. Crystallization may be used as a purification method during these steps.

In a similar fashion, monoclonal antibodies can be purified using crystallization from other sources including, *inter alia*, the following: insect cell culture; bacterial cell culture; plant parts including, *inter alia*, seeds, flowers, leaves and roots/tubers from transgenic plants, including, *inter alia*, transgenic maize, tobacco, potatoes and corn; and milk, serum, plasma, eggs, and other areas of transgenic animals, including, *inter alia*, transgenic cows, horses, pigs, chickens, goats and sheep.

Crystallization of antibodies may be carried out either directly from the cell extract before purification or after partial or complete purification, e.g., directly from milk or after clarification or

partial purification of the protein of interest at any stage of the process.

Method:

One liter of, e.g., Trastuzumab (at 22 mg/ml)  
5 is mixed in a beaker with one liter of a  
crystallization buffer containing 25% PEG400, 5%  
PEG8000, 100 mM Tris, pH 8.5, 10% propylene glycol, and  
0.1% Tween®80 (Sigma-Aldrich). The mixture is then  
incubated at room temperature overnight with stirring,  
10 using a overhead stirrer. The solution is then seeded  
with Trastuzumab crystals obtained from hanging drop or  
microbatches. 100 ml of propylene glycol is added and  
the mixture is stirred, until crystals are formed (for  
approximately 24 hours).

15 Example 42

Purification of Antibodies by Crystallization  
from Milk

Milk (purchased from a local farm and stored  
frozen at 70°C) is thawed at 37°C and de-fatted by  
20 centrifugation at 7000 x g for 15 minutes at 4°C. The  
cream layer is then punctured using a sharp pipette tip  
and the skim milk is decanted into a clean tube through  
the opening. Skim milk is then diluted with an equal  
volume of 250 mM EDTA. The milky appearance clears,  
25 indicating the destruction of micellar structures and  
aggregates. EDTA-clarified skim milk is dialyzed  
against PBS to remove EDTA. The clarified milk may  
then be spiked with monoclonal antibody solutions to a  
final concentration of approximately 5-10 mg/ml. The  
30 monoclonal antibody from the spiked milk is then  
crystallized using polyethylene glycol alone or in  
combination with salts, e.g., ammonium sulfate, or any  
of the crystallization conditions described earlier

using a variety of salts, buffer, organic solvents etc. This method may be used to purify antibodies from transgenic milk.

#### Example 43

5                   Crystallization of Monoclonal Antibodies from Transgenic Animals, Transgenic Animal Products, and Transgenic Plants

It will be understood by those of skill in the art that the method shown in Examples 41 and 42 may  
10 be use to purify monoclonal antibodies by crystallization, from transgenic animals (from cells, tissue extracts, etc.) transgenic animal products (e.g., eggs etc.) and from transgenic plants (plant cells and tissue extracts, etc.).

15                   In Examples 44 and 45, the purity and conformation of crystallized Rituximab was assessed by analyzing dissolved Rituximab on HPLC and SDS-PAGE, under reducing and non-reducing conditions.

#### Example 44

20                   Dissolved Rituximab from crystals obtained in Example 1 were analyzed on a 4-20% gradient SDS-PAGE gel, without the presence of  $\beta$ -mercaptoethanol. The omission of  $\beta$ -mercaptoethanol from the gel electrophoresis was to prevent the dissociation of the  
25 heavy and light chains, which are held together by a disulfide bond. Native Rituximab was analyzed under the same conditions as a control.

#### Results:

Both the native and dissolved Rituximab  
30 showed a single protein band under non-reducing conditions, with a molecular weight approximately equal

to 150 kD, the correct size for the whole Rituximab monoclonal antibody.

The protocol was repeated, using reducing conditions (using  $\beta$ -mercaptoethanol). Native Rituximab  
5 was analyzed under the same conditions as a control.

Results:

For both native and dissolved Rituximab under reducing conditions, the gel revealed two bands at about 50 kD and about 25 kD, the correct size for the  
10 monoclonal antibody heavy and light chains, respectively.

Example 45

Purity and Size Analysis of Rituximab  
15 Crystals by HPLC

The purity and size of native Rituximab and dissolved Rituximab from crystals obtained in Example 1 were analyzed with a size exclusion column (BIOSEP-SEC S3000, Phenomenex, Torrence, CA) on an HPLC (Shimadzu,  
20 LC-10AD) system with 100 mM potassium phosphate, pH 7.5 as running buffer and a constant flow rate of 0.5 ml/min. Both the native and dissolved Rituximab were eluted from the column as a single protein peak, indicating that the crystallization process did not  
25 alter the structural integrity of the Rituximab antibody.

Example 46

Dynamic Light Scattering Characterization of  
30 native (soluble) and dissolved Rituximab

Soluble Rituximab and dissolved Rituximab from Rituximab crystals, prepared as in Example 21, were dissolved in 25 mM Tris, pH 7.0, 150 mM sodium chloride and 0.1% Tween®80 (Sigma-Aldrich) (final

protein concentration equal to approximately 1 mg/ml), and analyzed on a PD2000 Dynamic Light Scattering detector with Precision/Acquire and Precision/Analyze (Precision Detectors, Franklin, MA). Native (soluble) Rituximab (10 mg/ml) was diluted 20-fold with deionized water for comparison.

Results:

The hydrodynamic radius for the soluble and dissolved Rituximab are identical, indicating that the crystallization process did not alter this characteristic of the Rituximab antibody. In addition, this example showed that no protein aggregation occurred during crystallization, as judged from measuring the hydrodynamic radius.

15 Example 47

Dynamic Light Scattering Characterization of native (soluble) and dissolved Trastuzumab

Trastuzumab crystals, prepared as in Example 31, were dissolved in 25 mM Tris, pH 7.0, 150 mM Sodium chloride and 0.1% Tween<sup>®</sup>80 (Sigma-Aldrich). The final protein concentration was adjusted to approximately 1 mg/ml. The dissolved Trastuzumab crystal was then analyzed on a PD2000 Dynamic Light Scattering detector with Precision/Acquire and Precision/Analyze (Precision Detectors, Franklin, MA). Native (soluble) Trastuzumab (22 mg/ml) was diluted 20-fold with deionized water for comparison.

Results:

The hydrodynamic radius and the molecular weight for the soluble and dissolved Trastuzumab were identical, indicating that the crystallization process did not alter this characteristic of the Trastuzumab antibody. In addition, this example showed that no

protein aggregation occurred during crystallization, as judged from measuring the hydrodynamic radius.

#### Example 48

The needle cluster Rituximab crystals, which  
5 were crystallized in Example 1 with 20% (w/v) PEG-1000, 100 mM imidazole, and 200 mM calcium acetate, pH 7.0, were determined to have a median of 72  $\mu$ m and a size range between 50-150  $\mu$ m, as characterized with a Coulter LS Particle Size Analyzer.

#### 10 Example 49

##### Peptide mapping comparison of native (soluble) and dissolved Rituximab

A 500  $\mu$ l aliquot of 10 mg/ml of Rituximab (soluble or crystals from Example 1) in 25 mM Tris, pH  
15 7.0 and 0.1% Tween®80 (Sigma-Aldrich) was mixed with 167  $\mu$ g of trypsin and incubated at 37°C for 24 hr. Each digested sample was filtered through a 0.22  $\mu$ m filter and a 200  $\mu$ l aliquot was loaded onto a C-8 reverse phase (Vydac, Hesperia, CA) HPLC column, which  
20 was equilibrated with water supplemented with 0.1% trifluoroacetic acid, on a Shimazu LC10AD system. The bound peptide was eluted with a 0-70% acetonitrile gradient over 70 min at 0.9 ml/min.

##### Results:

25 Similar profiles were obtained for soluble and redissolved Rituximab, indicating no change in conformation, structure or size of the Rituximab molecule due to the crystallization process.

#### Example 50

30 N-terminal sequencing of native (soluble) and

dissolved Rituximab and Trastuzumab:

In order to demonstrate that antibodies, e.g., Rituximab and Trastuzumab, do not suffer terminal degradation in the crystalline state, the following was performed on Rituximab and Trastuzumab crystals prepared according to Examples 21 and 31, respectively, and their soluble counterparts:

N-terminal sequencing was carried out using an Applied Biosystems, Inc. (ABI) 447A automatic protein sequencer. Each sample was loaded onto a glass fiber disc, which had been placed in the sequencer and pre-cycled once. Following the pre-cycling step, a number of cycles of Edman degradation were performed using a standard protein sequencing program from ABI. The results are reported as the major phenylthiohydantoin (PTH)-amino acid detected for each cycle. (Standard one-letter designations for the 20 commonly occurring amino acids are used to report the resulting sequences. They are: A = alanine; C = cysteine; D = aspartic acid; E = glutamic; F = phenylalanine; G = glycine; H = histidine; I = isoleucine; K = lysine; L = leucine; M = methionine; N = asparagine; P = proline; Q = glutamine; R = arginine; S = serine; T = threonine; V = valine; W = tryptophan; Y = tyrosine.)

Results:Trastuzumab:

Trastuzumab Form	Antibody Chain	N-terminal sequence
Crystalline	Heavy	E-V-Q-L-V-G-S
Crystalline	Light	D-I-Q-M-T-Q-S
Soluble	Heavy	E-V-Q-L-V-G-S



Soluble	Light	D-I-Q-M-T-Q-S
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Rituximab:

	Rituximab Form	Antibody Chain	N-terminal sequence
	Crystalline	Heavy	blocked
5	Crystalline	Light	Q-I-V-L-S-Q-S
	Soluble	Heavy	blocked
	Soluble	Light	Q-I-V-L-S-Q-S

The results show that the crystallization process does result in N-terminal amino acid degradation of the

10 Trastuzumab or Rituximab antibodies.

Example 51

PAS (Periodic acid-Schiff reagent) total carbohydrate staining of native (soluble) and dissolved Trastuzumab

15 Method:

2  $\mu$ g of a dialyzed sample of crystallized Trastuzumab, prepared according to Example 31, as well as its soluble counterpart (as supplied by the manufacturer), both reduced with 10%  $\beta$ -mercaptoethanol

20 and unreduced, were run on SDS-PAGE gels. The gels were transferred via Western blot to nitrocellulose membranes. Using the BioRad Immun-Blot staining kit, the carbohydrate moieties attached to the antibodies (and the heavy chains of the antibodies in the case of

25 the reduced samples) were stained and visualized. Specifically, the gels were fixed with 40% methanol, 7% acetic acid for 30 minutes, and washed 4 times in this solution, then left in overnight in fresh solution. The next day, gels were oxidized in 1% periodic acid,

3% acetic acid, and subsequently washed 10 times in double-distilled (dd) H<sub>2</sub>O, 10 minutes each, to remove periodic acid. Gels were incubated in the dark with Schiff's reagent for about one hour to develop stain  
5 and then scanned.

Results:

Both soluble and crystalline Trastuzumab appeared to be nearly identical on the gel, except for the light high-molecular weight band in the non-reduced  
10 sample of crystalline antibody. In the reduced sample, the carbohydrate was associated with the heavy chain of the antibody, as expected.

Example 52

PAS (Periodic acid-Schiff reagent) total  
15 carbohydrate staining of native (soluble) and  
dissolved Rituximab

Method:

This method was performed as in Example 51.

Results:

20 Soluble and crystallized samples of Rituximab appeared identical on the gel slab, with the carbohydrate associated with the heavy chain of the antibody in the reduced sample.

Example 53

25 N-linked oligosaccharide profiling of native  
(soluble) and dissolved Trastuzumab and  
Rituximab

N-linked Oligosaccharide Profiling was achieved using a BioRad N-linked Oligosaccharide  
30 Profiling Kit (catalog # 170-6501). Crystallized samples of Trastuzumab and Rituximab (from Examples 31 and 21, respectively) were washed, dissolved, and

dialyzed against ddH<sub>2</sub>O, and samples of soluble Trastuzumab and Rituximab (as supplied by manufacturer) were dialyzed against ddH<sub>2</sub>O. A 200 µg aliquot of each sample was mixed with equal volumes of releasing buffer and denatured using 1 µl 5% sodium dodecyl sulfate (SDS), 1.5 µl 10% β-ME, and 4 µl NP-40 (Tergitol at room temperature. Subsequently, 2 µl of PNGase (an enzyme that cleaves asparagine-linked oligosaccharides) was added to each sample, and samples were incubated overnight at 37°C. Protein was then precipitated with 3 volumes of cold ethanol, samples were spun, and supernatants (containing oligosaccharides) were recovered and lyophilized. Samples were reconstituted and fluorescently labeled with 8-aminonaphthalene-1,3,6-trisulfonic acid (ANTS) at 37°C overnight. Samples were again lyophilized and reconstituted in H<sub>2</sub>O and 2X sample buffer. Oligosaccharides were run in gel electrophoresis using N-linked oligosaccharide profiling gels and buffers from BioRad, and gels were visualized using a long-wave UV transilluminator.

#### N-linked oligosaccharides

N-linked oligosaccharides were cleaved from the antibodies and fluorescently stained using the BioRad N-linked Oligosaccharide Profiling Kit. The glucose ladder standard run to the left of the samples showed the relative positions of the oligosaccharides.

#### Results:

The bands for crystal and soluble Rituximab and Trastuzumab appeared to be identical, suggesting that crystallization causes no changes in the oligosaccharide groups associated with the starting material.

Example 54Monosaccharide Constitution of native  
(soluble) and dissolved Rituximab and  
Trastuzumab

5           The monosaccharide constituents of native  
(soluble) Rituximab and Trastuzumab were compared with  
those of reconstituted Rituximab and Trastuzumab  
crystals, as prepared in Examples 1 and 31,  
respectively, using a Bio-Rad Monosaccharide  
10 Composition Analysis kit (catalog # 170-6811).  
Trastuzumab and Rituximab crystals were washed three  
times in mother liquor and dialyzed overnight versus  
ddH<sub>2</sub>O. Reconstituted native (soluble) Trastuzumab and  
Rituximab were dialyzed versus ddH<sub>2</sub>O and used for  
15 comparison.

          The samples of native and dissolved  
antibodies were then analyzed for their sugar content.  
Each native and dissolved antibody was divided into  
three 50 µl aliquots containing 40 micrograms of  
20 antibody, one aliquot each for three hydrolysis  
reactions. For the neutral sugar analysis, the  
reaction took place in 2 N trifluoroacetic acid (TFA) at  
100°C over 5 hours. For amine sugars, the hydrolysis  
took place in 4N HCl at 100°C over 3 hours, and sialic  
25 acid hydrolysis was performed using 0.1 N TFA at 80°C  
for 1 hour. After hydrolysis, all samples were  
lyophilized, and the amine sugars were re-acetylated  
with acetic anhydride and a buffer, and lyophilized  
again. Samples were then reconstituted and  
30 fluorescently labeled with 2-aminoacridine (AMAC), then  
incubated at 45°C for 3.5 hours. Samples were  
lyophilized again, diluted, and mixed with 2X sample  
buffer for electrophoresis. Electrophoresis of samples

was accomplished using Bio-Rad Monosaccharide Composition Gels and buffers, and the results were visualized using a long-wave UV transilluminator.

Results:

5                   Rituximab: Native (soluble) and dissolved Rituximab had identical monosaccharide constituents. The neutral monosaccharides appearing on the gel were mannose, a small band of fucose, a small band of glucose (which could just be contamination, as it shows  
10 up in the blank) and a small band of galactose. N-glucosamine was the only band that appears after amine hydrolysis. The sialic acid hydrolysis yielded no bands.

                  Trastuzumab: Native (soluble) and dissolved  
15 Trastuzumab had identical monosaccharide constituents. The neutral monosaccharides appearing on the gel were mannose, a small band of fucose, and a small band of glucose (which could just be contamination, as it shows up in the blank). N-glucosamine was the only band that  
20 appeared after amine hydrolysis. The sialic acid hydrolysis yielded no bands.

                  The results for both Rituximab and Trastuzumab demonstrate that the crystallization process did not alter the monosaccharide content of the  
25 antibody.

Examples 55 and 56

Antibody Bioassays

                  Various monoclonal antibodies which recognize tumor-associated antigens, including, *inter alia*, those  
30 referred to herein, are widely used for cancer treatment. The cytotoxicity of an antibody on its antigen-bearing target cells can be characterized by

any one of three assays, e.g. direct cytotoxicity, complement dependent cytotoxicity (CDC), and Antibody-dependent cytotoxicity (ADCC). The target cells for Rituximab are the cells that overexpress CD-20 antigen on their surface, which include Raji, Daudi, JOK1 and WT100. The specific antigen for Trastuzumab is HER2 (human epidermal growth factor receptor 2 protein), which is overexpressed in human breast adenocarcinoma cell lines including SK-BR-3, BT474, and MCF/HER2.

10           1. Direct Cytotoxicity:

Direct cytotoxicity, as the name implies, measures the intrinsic toxic effect of an antibody on the target cell by co-incubating the target cells with different concentrations of antibody. Cell viability is counted after co-incubation with antibody.

2. Complement dependent cytotoxicity (CDC):

When an antibody binds to its cell surface antigen, it induces target cell destruction by activating the complement system (a series of interacting proteins that lyse cells and trigger local inflammatory reactions). This assay is carried out by co-incubating the fixed number of target cells with diluted human serum (as a source of complement system) and antibody. The cell viability is determined at the end of the incubation. Compared with the control plates (target cells plus antibody only), the cell death in the plates containing complement (human serum) is significantly elevated.

3. Antibody-dependent cytotoxicity (ADCC):

30           Similar to CDC, ADCC is one of the major mechanisms responsible for cytotoxicity of monoclonal antibodies. In contrast to CDC, the target cell destruction caused by ADCC is initiated by recruiting

immune cells, which specifically attack tumor cells, after an antibody binds to its specific antigen on the target cell. The ADCC assay is carried out by first seeding the wells/plates with fixed amount of target  
5 cells before co-incubated with antibody and effector immune cells (usually use the isolated peripheral blood mononuclear cells). The cell viability is determined at the end of co-incubation. Cell death is significantly increased with the presence of the  
10 effector immune cells, as compared with the control (target cell plus antibody only).

#### Example 55

##### Rituximab Crystals Induce Direct Cytotoxicity against the RAJI Lymphoma Cell Line

15 RAJI lymphoma cells (ATCC, Manassas, VA, ATCC # CCL 86) were cultured in growth media and diluted to  $0.5 \times 10^5$  cells/ml. A 100  $\mu$ l aliquot of that culture was transferred to one well of a 96-well plate and cultured in the presence of various concentrations of native  
20 (soluble) and dissolved Rituximab (from crystals prepared according to Example 21) for 3 days. The number of viable cells remaining after the three-day incubation was determined using CellTiter 96<sup>®</sup> Aqueous One Solution Cell Proliferation Assay (Promega Corp.,  
25 Madison, WI). (Figure 8).

##### Results:

Dissolved Rituximab induced Direct Cytotoxicity as capably as native Rituximab under identical conditions.

Example 56Rituximab Crystals Induce Complement-  
Dependent Cytotoxicity against RAJI Lymphoma  
Cells

5 RAJI lymphoma cells (ATCC, Manassas, VA) were  
cultured in growth media and diluted to  $0.5 \times 10^5$   
cells/ml. A 100  $\mu$ l aliquot of above culture was  
transferred to one well of a 96-well plate and cultured  
in the presence of 25  $\mu$ g/ml of native or dissolved  
10 Rituximab (from crystals prepared according to Example  
21) and various concentrations of human serum for 3  
days. The number of viable cells remaining after the  
three-day incubation was determined using CellTiter 96<sup>®</sup>  
Aqueous One Solution Cell Proliferation Assay (Promega  
15 Corp., Madison, WI). (Figure 9).

Results:

Dissolved Rituximab induced Complement-  
Dependent Cytotoxicity as capably as native Rituximab  
under identical conditions.

20 Example 57Cumulative Analysis Comparing Native  
(Soluble) and Crystalline Rituximab

The following table summarizes Examples 44-  
46, 49, 50, 54, 53, 55 and 56, respectively, comparing  
25 the properties of native (soluble) and crystalline  
Rituximab:

Analytical Methods	Soluble	Crystalline	Result
30 <u>SDS-PAGE</u> non-reducing conditions	Whole Ab MW = ~ 150 kD	Whole Ab MW = ~ 150 kD	Soluble and crystalline forms of Rituximab were



5

Analytical Methods	Soluble	Crystalline	Result
reducing conditions	H chain MW = ~ 50 kD L chain MW = ~ 25 kD	H chain MW = ~ 50 kD L chain MW = ~ 25 kD	identical. Crystallization did not alter the structural integrity of Rituximab.
<u>HPLC gel filtration</u>	Single peak	Single peak	Crystallization did not alter the structural integrity of Rituximab.
<u>Dynamic Light Scattering</u>	MW = ~ 150 kD	MW = ~ 150 kD	Crystallization did not alter the structural integrity of Rituximab or change the hydrodynamic radius.
<u>Peptide mapping</u>	Trypsin digest	Trypsin digest	Similar profiles were obtained for soluble and redissolved Rituximab, indicating no change in conformation, structure or size of the Rituximab molecule.
<u>N-terminal Sequencing of Antibody Light Chains</u>	Gln-Ile-Val-Leu-Ser-Gln-Ser	Gln-Ile-Val-Leu-Ser-Gln-Ser	Native (soluble) and dissolved Rituximab had identical N-

10

Analytical Methods	Soluble	Crystalline	Result
			terminal sequences, indicating no hydrolysis of amino acids from the N-terminal side.
<u>Monosaccharide Constitution</u>	Fucose, mannose, N-acetyl glucosamine, galactose	Fucose, mannose, N-acetyl glucosamine, galactose	Native (soluble) and dissolved crystalline Rituximab had identical monosaccharide constituents, indicating that no monosaccharides were cleaved from the monoclonal antibody during crystallization.
15 <u>Oligosaccharide Profiling</u>	Three bands Corresponding to G8, G9 and G10, corresponding to 8-, 9-, and 10-residue sugars.	Three bands Corresponding to G8, G9 and G10, corresponding to 8-, 9-, and 10-residue sugars.	Native (soluble) and dissolved crystalline Rituximab had identical oligosaccharide profiles, indicating that crystallization does not alter the oligosaccharide make-up of the antibody.
<u>Bioassays</u>  Direct Cytotoxicity	Yes	Yes	Native and dissolved Rituximab both induced each function. Thus,

Analytical Methods	Soluble	Crystalline	Result
Induced Complement Dependent Cytotoxicity	Yes	Yes	crystallization did not result in changes to immune functions.

### 5 Example 58

#### Secondary structure characterization by FTIR

The Fourier transform infrared (FTIR) spectra are collected on a Nicolet model 550 Magna series spectrometer as described by Dong et al. [Dong, A.,  
 10 Caughey, B., Caughey, W.S., Bhat, K.S. and Coe, J.E. *Biochemistry*, 1992; 31:9364-9370; Dong, A. Prestrelski, S.J., Allison, S.D. and Carpenter, J.F. *J.Pharm. Sci.*, 1995; 84: 415-424.]

For the solid samples, 1 to 2 mg of the  
 15 antibody or antibody fragment are lightly ground with 350 mg of KBr powder and filled into small cups used for diffuse reflectance accessory. The spectra are collected and then processed using Grams 32 (from Galactic Software) for the determination of relative  
 20 areas of the individual components of secondary structure using second derivative and curve-fitting program under amide I region (1600 -1700 cm<sup>-1</sup>).

The correlation coefficient is calculated using protein analysis software from Nicolet which  
 25 easily allows the determination of the correlation coefficient between the previously saved reference spectrum and that of the current protein spectrum (Garland, B, *FT-IR Studies of Protein Secondary Structure in Aqueous and Dried States*. Nicolet  
 30 application note # AN 9479). The second derivative

spectrum of the native aqueous protein is used as a reference spectrum and the dried crystals and lyophilized solid protein can be used as samples. The proteins will have an increasingly similar secondary conformational structure as the correlation coefficient approaches unity.

#### Results:

The correlation coefficient for a given crystalline monoclonal antibody in slurry form, or in dried crystalline form, when compared to soluble form (reference spectrum equals to one) is greater than 0.8 but less-than-or-equal-to 1.

#### Example 59

##### Soluble whole antibody sample preparation

For comparison to the Rituximab antibody crystals produced in Example 1, a sample of soluble whole antibody was prepared by dissolving (resuspending) whole antibody crystals to 20 mg/ml in 0.1% Tween®80 (Sigma-Aldrich), 150 mM sodium chloride and 25 mM Tris-HCl, pH 7.0 at 37°C. This method was used to dissolve Rituximab crystals for Examples 44, 45, 46, 49, 50, 52, 53, 54, 55 and 56. This method was also used to dissolve Trastuzumab crystals for Examples 47, 50, 51, 53 and 54. This method was also used to dissolve Infliximab crystals for Example 63.

#### Example 60

##### Crystallinity

The crystal integrity of the crystals and formulations and compositions thereof of this invention

may be measured by quantitative microscopic observations. In order to visualize whether the crystals maintained their shape after drying, dried crystals may be examined under an Olympus BX60  
5 microscope equipped with DXC-970MD 3CCD Color Video Camera with Camera Adapter (CMA D2) with Image ProPlus software. Samples of dried crystals can be covered with a glass coverslip, mounted and examined under 10X magnification, using an Olympus microscope with an  
10 Olympus UPLAN F1 objective lens 10X/0.30 PH1 (phase contrast).

#### Example 61

##### Trastuzumab Animal Models

Trastuzumab may be used in the treatment of  
15 breast cancer [Pietras R.J., Poen J.C., Gallardo, D., Wongvipat P.N., Lee H.J. and Slamon D.J., Cancer Res, vol. 59, pp. 1347-55 (1999); Baselga, J., Norton L., Albanell J., Kim Y.M., Mendelsohn J., Cancer Research, vol. 58, pp. 2825-31 (1998)].

##### 20 Procedure of Tumor Formation in Nude Mice:

Human breast cancer SK-BR3 or BT-474 cells (American Type Culture Collection (ATCC) (Manassas, Virginia, USA)) were cultured in BRMI 1640 medium supplemented with 10% fetal bovine serum (FBS), 2 mM  
25 glutamine and 1% penicillin G/streptomycin/fungizone solution. After a few cell passages, the human breast cancer cells were inoculated subcutaneously (s.c.) ( $5 \times 10^7$  cells/animal) in the hind thighs of 3-month-old female athymic mice.

30 Prior to inoculation, mice were primed for 10-14 days with  $17\beta$ -estradiol applied s.c. in a

biodegradable carrier-binder (1.7 mg of estradiol per pellet) to promote growth of the estrogen-dependent breast cancer cells. Tumor nodules were monitored by measuring their dimensions (in mm). Five to six  
5 animals were included in each treatment group. The animals were randomly chosen with respect to body weight and tumor nodule size at the start of each treatment. Antibody treatment was initiated when tumors grew to more than 50 mm<sup>3</sup> in size in one set of  
10 animals or to more than 350 mm<sup>3</sup> in size in a second set. Monoclonal antibodies and control solutions were administered by intra-peritoneal (i.p.) injection. Recombinant human (rhu) Mab HER-2 antibody (Trastuzumab) was given at a dose of 5 or 10 mg/kg  
15 animal body weight in three doses at 4-day intervals (over 12 days). Control injections were of human IgG1 (5 or 10 mg/kg), also given i.p., using the same administration protocol. Mice were then sacrificed for pathological examination.

20 The crystalline and soluble Trastuzumab eradicated most or all of the tumors formed by injecting BT 474 cells into mice, when compared to controls consisting of saline (which was used as the cell delivery vehicle) or non-specific IgG, clearly  
25 indicating that the crystalline Trastuzumab is efficacious in mice animal models for breast cancer.

Trastuzumab Pharmacokinetics (PK) Studies in Mice:

For PK studies of Trastuzumab, Trastuzumab  
30 was inoculated s.c. or i.v. into Balb/C Mice. 25 to 30 Balb/C mice of the same sex, weighing between 20 and 25 grams each, were used for the study. On day 1, mice

were weighed and then given a single subcutaneous or intravenous injection of Trastuzumab (30 mg Trastuzumab per kg mouse body weight). The concentration of the Trastuzumab solution/suspension was adjusted so that  
5 the prescribed dose was administered in a volume of 5 ml per Kg mouse body weight. At 0.5, 1, 2, 4, 6, 8, 24 and 48 hours after dosing, blood samples were obtained from 3 of the mice. Three mice at each time point were anesthetized and as much blood as possible was drawn  
10 from the heart (terminal bleed) and transferred to Microtainer serum-separator tubes. About 300 µl of blood were collected from each mouse. The collected blood was allowed to clot and the tubes were centrifuged. The blood cells were removed, and the  
15 supernatant (serum) was decanted into cryo-vials and frozen at -70°C. Subsequently, Trastuzumab levels were determined by ELISA (see protocol below), and the results were plotted as the number of µg Trastuzumab per ml serum over time post-inoculation.

20                   Trastuzumab ELISA Protocol:

Wells of 96-well high binding polystyrene plates from Sigma (Costar brand) were coated with 10 µg/ml anti-human antibody from Pierce (50 µl per well) at 4°C overnight. The anti-human antibody was removed  
25 and the plates were washed three times with a buffer containing 50 mM Tris, pH 8.0, 0.138 M sodium chloride, 0.0027 M potassium chloride and 0.05% Tween®20 (Sigma-Aldrich) (TBST). Each well was then blocked with 200 µl of 3% non-fat dry milk (Sigma) in TBST for 2 hours  
30 at room temperature. The plates were emptied and washed 3 times with TBST. Serum samples containing Trastuzumab were diluted in non-fat dry milk in TBST.

A 100  $\mu$ l aliquot was added to each appropriate well. A 100  $\mu$ l aliquot of control sample (either saline or IgG) was added to the appropriate wells and the plates were incubated for 2 hours at room temperature. The plates  
5 were emptied and washed 3 times with TBST. A 100  $\mu$ l aliquot of Horseradish peroxidase (Sigma) (a 1/25,000 dilution) conjugated anti-human antibody (in non-fat dry milk in TBST) was then added to each well and the plates were incubated for 1 hour at room temperature.  
10 The plates were then emptied and washed 3 times with TBST. 100  $\mu$ l of 3,3', 5,5'-Tetra methyl-benzidin (TMB) substrate (Sigma) was added to each well and the plates were incubated in the dark for 30 minute at room temperature in order to allow for the color reaction to  
15 proceed. The color reaction was stopped by adding 100  $\mu$ l 1 N sulfuric acid to each well. The absorbance was read at a wavelength of 450 nm ( $OD_{450}$ ) on an automatic Microplate reader. The  $OD_{450}$  values, which corresponded to the amount of Trastuzumab in the blood sample  
20 tested, were then plotted. The resulting plot is shown in Figure 15.

#### Results:

The crystalline Trastuzumab injected i.v. entered the bloodstream immediately (it had reached  
25 its approximate maximum serum level at the first time point = 30 minutes) and maintained its serum concentration for approximately 480 minutes before the serum levels started to drop. See Figure 15. The crystalline Trastuzumab that was injected s.c. took  
30 longer to enter the bloodstream (it reached its approximate maximum serum level in approximately 480 minutes) than the Trastuzumab administered i.v.



However, the approximate maximum serum levels of the Trastuzumab administered s.c. were maintained until at least the last time point = 48 hours, indicating that Trastuzumab crystals administered s.c. may  
5 advantageously maintain and control high serum levels of the antibody over extended periods of time. See Figure 15.

#### Example 62

##### Rituximab Animal Models

10 Rituximab may be used for the treatment of non-Hodgkins lymphoma [Bertolini, F., Fusetti, L., Mancuso, P., Gobbi, A., Corsini, C., Ferrucci, P.F., Blood, volume 96, pp. 282-87 (2000)].

##### Procedure of Tumor Formation in Nude Mice:

15 A model of high-grade human grade non-Hodgkins lymphoma was generated by injecting 6- to 8-week-old NOD/SCID mice intraperitoneal (i.p.) with  $10 \times 10^6$  Raji cells (ATCC). The mice were evaluated for tumor growth every other day. Tumor volume was  
20 measured with calipers, and the formula, width x length x 0.52, was applied to approximate the volume of the spheroid tumors. The chimeric anti-CD20 monoclonal antibody Rituximab, in its crystalline form, as prepared in Example 21, at concentrations ranging from  
25 25-75 mg/ml, was given intraperitoneal (i.p.) to mice on days 3, 5, and 7. Control mice received i.p. or subcutaneous (s.c.) injections of phosphate-buffered saline (PBS). Tumor-bearing mice were killed by carbon dioxide asphyxiation, and tumor engraftment was  
30 confirmed by histologic, immunohistochemistry (IHC), and flow cytometry (FC) studies.

Example 63Infliximab Animal Models

Infliximab has been used in the treatment of arthritis [Kim, S.H., Evans, C.H., Kim, S., Oligino, T., Ghivizzani, S.C. and Robbins, P.D., Arthritis Res., vol. 2, pp. 293-302 (2000); Yoshino, S., The Journal of Immunology, vol. 160, pp. 3067-71 (1998); Malfait, A.M., Williams, R.O., Malik, A.S., Maini, R.N. and Feldmann, M., Arthritis Rheum., vol. 44, pp. 1215-24 (2001)].

Procedure of Tumor Formation in Nude Mice:

Male DBA/1 lacJ (H-2q) mice, aged 7-8 weeks, were purchased from The Jackson Laboratory (Bar Harbor, ME, USA). The mice were immunized intradermally (i.d.) at the base of tail with 100 µg of bovine type II collagen. On day 21 after immunization, the mice received a booster injection (i.d.) of 100 µg type II collagen in Freund's incomplete adjuvant. For the synchronous onset of arthritis, 40 µg of lipopolysaccharide (Sigma, St Louis, MO, USA) was injected i.p. on day 28.

At the onset of clinical arthritis, mice were treated for 4 weeks with anti-tumor necrosis factor (anti-TNF; Infliximab). Two milligrams of Infliximab crystals, as prepared in Example 37, were dissolved in 0.5 ml of PBS and injected i.p. daily. For treatment controls, 0.5 ml of PBS only and 0.5 ml of PBS containing 2 mg of nonspecific IgG were given to mice. After 4 weeks of treatment, mice were killed and hind paws were assessed histologically for joint damage.

Example 64

Stability of Rituximab in the Crystalline Form

Assay 1:

A 0.35 ml aliquot of Rituximab (at 10 mg/ml) crystal slurry, as prepared in Example 21, was stored in mother liquor at room temperature. 5 µl aliquots were removed at different time points over a one-month period. The integrity of the antibody was then analyzed on a non-reducing 4-20% Tris-glycine gradient gel. A single protein band was observed on the gel after Coomassie blue stain.

Assay 2:

A 100 mg/ml aliquot of Rituximab crystal slurry, as prepared in Example 21, was prepared by centrifuging 1.1 ml of crystal slurry and pellet was resuspended in 50 µl buffer containing 150 mM Sodium chloride, 25 mM sodium citrate, 0.09% Tween®80 (Sigma-Aldrich), pH 6.5. The suspended slurry was then stored at room temperature for a month before being analyzed on a non-reducing 4-20% Tris-glycine gradient gel. A single protein band was observed on the gel. (Figure 12).

Example 65

Stability of Crystalline and Native (Soluble) Trastuzumab in the Presence of Organic Solvents

Crystalline Trastuzumab:

Trastuzumab was crystallized as described in Example 31. A 50 µl aliquot of a slurry of Trastuzumab crystals (with a protein concentration of 22 mg/ml) was

centrifuged in order to remove the mother liquor. The supernatant (mother liquor) was discarded. The Trastuzumab crystals were resuspended in 200  $\mu$ l of acetone. The crystal/acetone mixture was incubated for 3 hours at 4°C. The acetone was removed by centrifugation and the Trastuzumab crystals were dissolved in 100  $\mu$ l of a solution containing 50 mM Tris, pH 7.0, 100 mM sodium chloride. The dissolved (soluble) Trastuzumab was then analyzed by Size-exclusion chromatography on HPLC (SEC-HPLC), using a Phenomenex 2000 SEC column, a buffer consisting of 50 mM Tris, pH 7.0 and 100 mM Sodium chloride, and a flow rate of 0.5 ml/minute.

#### Native Trastuzumab:

A 50  $\mu$ l aliquot of soluble Trastuzumab (as supplied by manufacturer) at 22 mg/ml was added to 200  $\mu$ l acetone. The crystal/acetone mixture was incubated for 20 minutes at 4°C. The acetone was removed by centrifugation and the soluble Trastuzumab was analyzed by SEC-HPLC, using a Phenomenex 2000 SEC column, a buffer consisting of 50 mM Tris, pH 7.0 and 100 mM Sodium chloride, and a flow rate of 0.5 ml/minute.

#### Results:

The dissolved Trastuzumab remained whole and maintained its native structure (Figure 13), while the native/soluble Trastuzumab precipitated after the acetone treatment (Figure 14), demonstrating a loss of the structural integrity of the native Trastuzumab.

This example demonstrates the increased stability of Trastuzumab in the crystalline state according to this invention, as compared with the stability of the native Trastuzumab antibody.

As demonstrated herein, crystallization of monoclonal antibodies including, *inter alia*, Trastuzumab, results in increased stability and the preservation of the structural integrity of the antibodies. Stability in organic solvents is very useful when performing controlled dissolutions using various polymers, e.g., polylactic-co-glycolic acid ("PLGA"), for generating PLGA-encapsulated microspheres. Such processes cannot be performed if the antibodies or antibody fragments to be encapsulated become denatured in organic solvents.

Similar results have been obtained when Trastuzumab was exposed to other organic solvents, such as acetonitrile, ethanol, isopropyl alcohol (IPA) and 2-methyl-2,4-pentanediol ("MPD").

#### Example 66

##### Stability of Formulated Crystalline and Native (Soluble) Trastuzumab

##### Crystalline Trastuzumab:

Trastuzumab crystals were prepared as in Example 31.

50  $\mu$ l of a slurry containing crystals of Trastuzumab (at 22 mg/ml) were centrifuged, and the supernatant was discarded. The Trastuzumab crystals were resuspended in either buffer no. 1 (5% PEG 3350, 25% ethanol, 0.1 % Tween<sup>®</sup>80 (Sigma-Aldrich), 50 mM trehalose, 50 mM sodium phosphate, pH 7.6) or buffer no. 2 (25% PEG 3350, 5% alcohol (either ethanol or isopropanol), 0.1% Tween<sup>®</sup>80 (Sigma-Aldrich), 50 mM trehalose, 100 mM Tris, pH 8.0). The crystal/buffer mixture was incubated at 4°C for 16 days, after which

the buffer supernatant was removed by centrifugation, and the crystals were dissolved in 100 µl of 50 mM Tris, pH 7.0, 100 mM sodium chloride. The Trastuzumab was subsequently analyzed by SEC-HPLC (using a  
5 Phenomenex 2000 SEC column, a buffer consisting of 50 mM Tris, pH 7.0, 100 mM sodium chloride, and a flow rate of 0.5 ml/minute).

Native Trastuzumab:

50 µl of soluble Trastuzumab (as supplied by  
10 manufacturer) (at 22 mg/ml) was added to either buffer no. 1 or no. 2. The crystal/buffer mixture was incubated at 4°C for 2 hours, after which the buffer supernatant was separated from the precipitate, and the precipitate was analyzed for the presence of native  
15 Trastuzumab by SEC-HPLC (using the same conditions as above).

Results:

The dissolved crystalline Trastuzumab remained whole and maintained its native structure  
20 after being incubated with buffer 1 or 2, while the native/soluble Trastuzumab precipitated after being incubated with buffer 1 or 2, demonstrating a loss of the structural integrity of the native (soluble) Trastuzumab.

25 Example 67

Formulations Using Sucrose Acetate  
Isobutyrate (SAIB)

Formulations that use sucrose acetate isobutyrate (SAIB) relate to a parenteral liquid non-  
30 polymeric drug delivery system. This system comprises an antibody crystal or antibody fragment crystal

suspended in sucrose acetate isobutyrate (SAIB) and a "plasticizing" solvent, e.g., lanolin, mineral oil, ethanol, that is injected as a liquid [S.A. Sullivan, R.M. Gilley, J.W. Gibson and A.J. Tipton, 5 Pharmaceutical Research, vol. 14, p. 291 (1997)]. Following injection, the viscosity of the solution increases. The resulting high viscosity matrix is adhesive, biodegradable and biocompatible. The antibody is released in a controlled manner from the 10 matrix.

Trastuzumab Formulation Using Sucrose Acetate Isobutyrate (SAIB)

The SAIB formulation system outlined above was performed using the antibody Trastuzumab.

15 Method:

The mother liquor was removed from a 100 µl aliquot of a crystal slurry containing Trastuzumab crystals. Then, the crystals were washed with a solution of 90% SAIB in ethanol (ethanol acts as a 20 plasticizer in this example). At various time increments, 10 µl aliquot were removed from the SAIB/ethanol solution and suspended in 100 µl of 50 mM Tris, pH 7.0, 100 mM Sodium chloride. After a 10 minute incubation at room temperature, the crystals 25 were dissolved and the resulting material was analyzed on SEC-HPLC to determine the structural integrity of the Trastuzumab.

Results:

The monoclonal antibody remained stable under 30 the above conditions. Therefore, the SAIB formulation was shown to be suitable for use as a vehicle to deliver the Trastuzumab antibody subcutaneously.

It will be appreciated by those of skill in the art that the SAIB formulation according to this invention may be used as a controlled-delivery system for any crystal or crystal formulations or compositions of antibodies or fragments thereof, according to this invention.

#### Example 68

##### Stability of Crystalline Rituximab

For crystallization, 100  $\mu$ l of Rituximab in 9 mg/ml of sodium chloride, 7.35 mg/ml sodium citrate dihydrate, 0.7 mg/ml polysorbate 80, pH 6.5 was mixed with 200  $\mu$ l of crystallization buffer containing 0.1 M Hepes, pH 7.7, 12% PEG 400, 1.17 M sodium sulfate. The tube was then seeded with Rituximab crystals obtained from a microbatch and supplemented with 10  $\mu$ l of 1.5 M sodium sulfate before incubated at room temperature overnight. The stability of the crystals in ethanol or PEG or a combination of both was determined after taking 20  $\mu$ l of above crystal slurry and separating the crystals by centrifugation. The crystals were then suspended in 200  $\mu$ l of 25 mM Tris, pH 7.0, with or without the presence of 10% ethanol and 10% PEG 3350 at 37°C. Samples were taken at up to 24 hr for the presence of dissolved protein in the supernatant. The protein was determined by BioRad Protein Assay Kit I (BioRad Laboratories, Catalogue No. 500-0001).

##### Results:

The results indicate that the crystals were readily soluble in 25 mM Tris, pH 7.0, and 10% ethanol. The solution containing 10% PEG alone precipitated the



protein without maintaining the crystallinity. However, a combination of both ethanol and PEG maintained crystallinity and did not dissolve the crystals over a period of 1500 min (Figure 10). It is possible to use a combination of both ethanol and PEG formulation for the Rituximab crystals, since both of the reagents are pharmaceutically acceptable.

#### Example 69

##### Stability of Crystalline Trastuzumab

Trastuzumab was crystallized as described below. In brief, 200 ml of Trastuzumab (22 mg/ml in original formulation, as provided by the manufacturer) with equal volume of crystallization buffer containing 25% PEG 400, 5% PEG 8000, 10% propylene glycol, 100 mM Tris, pH 8.0, 0.1% Tween®80 (Sigma-Aldrich) and incubated overnight at room temperature. The tube was then seeded and supplemented with 20 µl propylene glycol. The stability of the crystals in ethanol or PEG or a combination of both was determined after taking 80 µl of above crystal slurry and separating the crystals by centrifugation. The crystals were then suspended in 400 µl of 25 mM Tris, pH 7.0 with or without the presence of 10% ethanol and 10% PEG 3350 at 37°C. Samples were taken at up to 140 hours for the presence of dissolved protein in the supernatant. The protein was determined by BioRad Protein Assay Kit I (BioRad Laboratories - Catalogue No. 500-0001).

##### Results:

The results indicate that the crystals were readily soluble in 25 mM Tris, pH 7.0 and 10% ethanol.

The solution containing 10% PEG alone precipitated the protein without maintaining the crystallinity. However, a combination of both ethanol and PEG maintained crystallinity and did not dissolve the crystals over a period of 8400 minutes (Figure 11). It is possible to use a combination of both ethanol and PEG formulation for the Trastuzumab crystals, since both of these reagents are pharmaceutically acceptable.

#### Example 70

10           Preparation of whole antibody  
              crystals using polyethylene oxide (PEO) as  
              excipient

In order to enhance the stability of whole antibody crystals prepared according to this invention during drying and storage, the crystals may be formulated with excipients. Whole antibody crystals according to this invention may be formulated using 0.1% polyethylene oxide in water as follows. The crystals are separated from the mother liquor by centrifugation at 1000 rpm in a Beckman GS-6R bench top centrifuge equipped with swinging bucket rotor. Next, the crystals are suspended in 0.1% polyethylene oxide for 3 hrs (Sigma Chemical Co., St. Louis, MO) and then separated by centrifugation.

#### 25   Example 71

Preparation of whole antibody crystals  
              using sucrose as excipient

Whole antibody crystals according to this invention may be formulated in the slurry form in the presence of mother liquor before drying. Sucrose

- (Sigma Chemical Co., St. Louis, MO) is added to whole antibody crystals in mother liquor as an excipient. Sufficient sucrose is added to whole antibody crystals to reach a final sucrose concentration of 10% (w/v).
- 5 The resulting suspension is then tumbled at room temperature for 3 hr. After treatment with sucrose, the crystals are separated from the liquid by centrifugation, as described in Example 70.

Example 72

- 10           Formulation of whole antibody crystals  
              using trehalose as excipient

- Whole antibody crystals according to this invention may be formulated as in Example 71, by adding trehalose instead of sucrose, (Sigma Chemical Co., St.
- 15 Louis, MO), to a final concentration of 10% (w/v) in mother liquor. The resulting suspension is then tumbled at room temperature for 3 hr and the crystals are separated from the liquid by centrifugation, as described in Example 70.

20   Example 73

Formulation of whole antibody  
crystals using methoxypolyethylene glycol  
(MOPEG) as excipient

- Whole antibody crystals are formulated as in
- 25 Example 71, by adding methoxypolyethylene glycol (Sigma Chemical Co., St. Louis, MO), instead of sucrose, to a final concentration of 10% (w/v) in mother liquor and separating after 3 hrs by centrifugation, as in Example 70.

Example 74Methods of drying crystal formulationsMethod 1. N<sub>2</sub> Gas Drying at Room Temperature

5           Crystals as prepared in one of Examples 1-3  
are separated from the mother liquor containing  
excipient by centrifugation at 1000 rpm in a Beckman  
GS-6R bench top centrifuge equipped with swinging  
bucket rotor in a 50 ml Fisher brand disposable  
10 centrifuge tube (Polypropylene). The crystals are then  
dried by passing a stream of nitrogen at approximately  
10 psi pressure into the tube overnight.

Method 2. Vacuum Oven Drying

15           Crystals, as prepared in one of Examples 1-3,  
are first separated from the mother liquor/excipient  
solution using centrifugation at 1000 rpm in a Beckman  
GS-6R bench top centrifuge equipped with swinging  
bucket rotor in a 50 ml Fisher brand Disposable  
polypropylene centrifuge tube. The wet crystals are  
20 then placed in a vacuum oven at 25 in Hg (VWR  
Scientific Products) at room temperature and dried for  
at least 12 hours.

Method 3. Lyophilization

25           Crystals, as prepared in one of Examples 1-3,  
are first separated from the mother liquor/excipient  
solution using centrifugation at 1000 rpm in a Beckman  
GS-6R bench top centrifuge equipped with swinging  
bucket rotor in a 50 ml Fisher brand Disposable  
polypropylene centrifuge tube. The wet crystals are  
30 then freeze dried using a Virtis Lyophilizer Model 24

in semi-stoppered vials. The shelf temperature is slowly reduced to -40 °C during the freezing step. This temperature is then held for 16 hrs. Secondary drying may then be then carried out for another 8 hrs.

5                    Method 4. Organic Solvent and Air Drying

Crystals, as prepared in one of Examples 1-3, are first separated from the mother liquor/excipient solution using centrifugation at 1000 rpm in a Beckman GS-6R bench top centrifuge equipped with swinging  
10 bucket rotor in a 50 ml Fisher brand Disposable polypropylene centrifuge tube. The crystals are then suspended in an organic solvent like ethanol or isopropanol or ethyl acetate or other suitable solvents, and centrifuged. The supernatant is then  
15 decanted and air dried at room temperature in the fume hood for two days.

Method 5. Air Drying at Room Temperature

Crystals, as prepared in one of Examples 1-3, are separated from the mother liquor containing  
20 excipient by centrifugation at 1000 rpm in a Beckman GS-6R bench top centrifuge equipped with swinging bucket rotor in a 50 ml Fisher brand Disposable centrifuge tube (Polypropylene). Subsequently, the crystals are then allowed to air dry in the fume hood  
25 for two days.

Method 6. Spray Drying

Crystals, as prepared in one of Examples 1-3, are spray dried using a Buchi Mini Spray Dryer Model B-  
30 191. The slurry of crystals at a concentration of 30

to 50 mg/ml is used for spray drying.

Example 75

Crosslinking of Antibody or Antibody Fragment  
Crystals and Formulations or Compositions  
5 Thereof

Crosslinking of crystals of a whole antibody, or crystals of an antibody fragment is carried out by incubating the crystal at a pH, such that the crosslinker is highly active and the crystalline nature  
10 of the antibody is preserved. Crosslinking is carried out either at ambient temperature or at 4°C with tumbling or stirring. After 24 hrs, the slurry is centrifuged at 3000 rpm and the supernatant is discarded. The excess (or un-reacted) crosslinker is  
15 inactivated with an appropriate buffer salt like Tris or glycine. The pellet is then washed with the mother liquor or appropriate pharmaceutically-acceptable buffer to remove the excess (or un-reacted) crosslinker. The crosslinking conditions may change  
20 depending on the type of crosslinker used but the ultimate goal is to maintain the crystalline state of the antibody under the conditions used.

It will be understood that antibody or antibody fragment crystals may be crosslinked using any  
25 suitable crosslinking reagent including, *inter alia*, Dimethyl 3, 3'-dithiobispropionimide.HCl (DTBP), Dithiobis (succinimidylpropionate) (DSP), Bis maleimido- hexane (BMH), Bis[Sulfosuccinimidyl]suberate (BS), 1,5-Difluoro-2,4-  
30 dinitrobenzene (DFDNB), Dimethylsuberimide.2HCl (DMS), Disuccinimidyl glutarate (DSG),

Disulfosuccinimidyl tartarate (Sulfo-DST), 1-Ethyl-3-[3-Dimethylaminopropyl]carbodiimide hydrochloride (EDC), Ethylene glycolbis[sulfosuccinimidylsuccinate] (Sulfo-EGS), N-[g-maleimidobutyryloxy]succinimide ester (GMBS), N-hydroxysulfosuccinimidyl-4-azidobenzoate (Sulfo-HSAB), Sulfosuccinimidyl-6-[a-methyl-a-(2-pyridyldithio)toluamido] hexanoate (Sulfo-LC-SMPT), Bis-[b-(4-azidosalicylamido) ethyl]disulfide (BASED) and glutaraldehyde (GA).

10 Example 76

Dissolution of Disulfide Bond-Containing  
Crosslinked Monoclonal Antibody Crystals

A 200 mM solution of cysteine is prepared by dissolving 242 mg of cysteine in 10 ml of 10 mM Tris HCl buffer, pH 7, containing 10 mM calcium chloride and 20% MPD. A 200 ml aliquot of a slurry of crosslinked monoclonal antibody crystals prepared according to this invention is taken and centrifuged at 3000 rpm for 5 minutes and the supernatant is discarded. The pellet is suspended in 200 ml of cysteine-containing Tris buffer. Another 200 ml of monoclonal antibody crystals is taken and centrifuged at 3000 rpm for 5 minutes and the supernatant is discarded. The pellet is then suspended in 200 ml of Tris buffer without any cysteine. All samples are incubated at 37°C for 1 hour and monitored for dissolution in 32 mM NaOH (direct visual and microscopic observation).

After incubation for 1 hour at 37°C, the DTBP sample is fully soluble in the presence of cysteine and insoluble in its absence. The DSP sample is barely

soluble in the presence of cysteine and insoluble in its absence.

Example 77

5                   Characterization of pH Solubility of  
                  Crosslinked Whole Antibody Crystals  
                  at 37°C

The solubility of various monoclonal antibody crystals prepared and crosslinked according to this invention with Dimethyl 3, 3'-  
10 dithiobispropionimidate.HCl (DTBP), Dithiobis (succinimidylpropionate) (DSP), Bis maleimido hexane (BMH), Bis[Sulfosuccinimidyl]suberate (BS), 1,5-Difluoro-2,4-dinitrobenzene (DFDNB), Dimethylsuberimidate.2HCl (DMS), Disuccinimidyl  
15 glutarate (DSG), Disulfosuccinimidyl tartarate (Sulfo-DST), 1-Ethyl-3-[3-Dimethylaminopropyl]carbodiimide hydrochloride (EDC), Ethylene glycolbis[sulfosuccinimidylsuccinate] (Sulfo-EGS), N-[g-maleimidobutyryloxy]succinimide ester (GMBS), N-  
20 hydroxysulfosuccinimidyl-4-azidobenzoate (Sulfo-HSAB), Sulfosuccinimidyl-6-[a-methyl-a-(2-pyridyldithio)toluamido] hexanoate (Sulfo-LC-SMPT), Bis-[b-(4-azidosalicylamido) ethyl]disulfide (BASED) and glutaraldehyde (GA) may be assayed.

25                   In 1.5 ml Eppendorf tubes, samples of uncrosslinked monoclonal antibody crystals and crosslinked monoclonal antibody crystal slurry, equal to 2.8 mg enzyme, may be microfuged at 3000 rpm for 5 minutes until the supernatant liquid is removed. Two  
30 pHs are tested: a) pH 7.4 and b) pH 2.0.

For pH 7.4, a 200 ml aliquot of PBS buffer



(0.01 M phosphate, 0.0027 M potassium chloride, 0.137 M sodium chloride, pH 7.4) is added to each sample, bringing the concentration of monoclonal antibody to 14 mg/ml. The samples may then incubated at 37°C for 24  
5 hours.

For pH 2.0, a 200 ml aliquot of glycine/HCl buffer pH 2.0 is added to each sample, bringing the concentration of monoclonal antibody to 14 mg/ml. The samples are incubated at 37°C for 5 hours. Initially,  
10 the samples are treated with 10 mM glycine/HCl buffer, pH 2.0 containing 10 mM calcium chloride and 20% MPD overnight at 25°C with tumbling; then proceeding with glycine/HCl buffer alone.

Samples may be studied for dissolution by  
15 centrifuging the samples at 14,000 rpm for 5 minutes after 5 hours and 24 hours. After centrifugation, the supernatant is passed through a 0.22 mm filter. The protein (antibody) content of the supernatant is then estimated then by removing 2 µl of the aliquot and  
20 placing it in 798 ml of deionized water. A 200 µl aliquot of Bio-Rad Protein assay reagent is added to this sample and is incubated at ambient temperature for 5 minutes and measured at 595 nm wavelength (Bio-Rad micro protein assay by Bradford's method). As a  
25 standard, 0-20 µg bovine IgG (Sigma) is used.

While we have described a number of embodiments of this invention, it is apparent that our basic examples may be altered to provide other embodiments which utilize the products and processes of  
30 this invention. Therefore, it will be appreciated that the scope of this invention is to be defined by the

appended claims rather than by the specific embodiments which have been represented by way of example.

CLAIMS

We claim:

1. A crystal of a whole antibody.
2. A crystal of a single-chain Fv fragment of an antibody.
3. A crystal of an Fab fragment of an antibody.
4. The crystal according to any one of claims 1, 2 or 3, wherein said whole antibody or single-chain Fv fragment or Fab fragment is characterized by  $\beta$ -sheet structural content of the antibody or fragment, as indicated by a correlation spectra as compared to the soluble counterpart of said antibody or antibody fragment, determined by FTIR, that is between about 0.8 and about 1.0.
5. The crystal according to any one of claims 1, 2 or 3, wherein said whole antibody or single-chain Fv fragment or Fab fragment is a therapeutic antibody or antibody fragment.
6. The crystal according to any one of claims 1, 2 or 3, wherein said whole antibody or single-chain Fv fragment or Fab fragment is a polyclonal antibody, or fragment thereof, or a monoclonal antibody, or fragment thereof.
7. The crystal according to any one of

claims 1, 2 or 3, wherein said crystal is a carrier-free pharmaceutical controlled release crystal.

8. The crystal according to any one of claims 1, 2 or 3, wherein said antibody is selected from the group consisting of: chimeric antibodies, humanized antibodies, non-glycosylated antibodies, bispecific antibodies, human antibodies and mouse antibodies.

9. The crystal according to any one of claims 1, 2 or 3, wherein said antibody is selected from the group consisting of: IgG, IgM, IgA, IgD, IgE, and serum IgA antibodies.

10. The crystal according to claim 9, wherein said antibody is selected from the group consisting of: IgG1, IgG2, IgG3 and IgG4, IgM1 and IgM2, and IgA1 and IgA2 antibodies.

11. The crystal according to any one of claims 1, 2 or 3, wherein said whole antibody or single-chain Fv antibody fragment or Fab antibody fragment has a greater half life in vivo than the soluble counterpart of said antibody or antibody fragment.

12. The crystal according to any one of claims 1, 2 or 3, wherein said antibody is an anti-idiotypic antibody.

13. The crystal according to any one of

claims 1, 2 or 3, wherein the antibody is selected from the group consisting of: Rituximab, Infliximab and Trastuzumab.

14. The crystal according to any one of claims 1, 2 or 3, wherein the antibody is selected from the group consisting of:

Abciximab,  
Palivizumab,  
Murumonab-CD3,  
Gemtuzumab,  
Trastuzumab,  
Basiliximab,  
Daclizumab,  
Etanercept, and  
Ibritumomab tiuxetan.

15. The crystal according to any one of claims 1, 2 or 3, wherein said antibody is selected from the group consisting of: anti-TNF antibodies, anti-CD3 antibodies, anti-CD20 antibodies, anti-CD25 antibodies, anti-CD33 antibodies, anti-CD40 antibodies, anti-HER2 antibodies, anti-HBV antibodies, anti-HAV antibodies, anti-HCV antibodies, anti-GPIIb/IIIa receptor antibodies, anti-RSV antibodies, anti-HIV antibodies, anti-HSV antibodies and anti-EBV antibodies.

16. The crystal according to any one of claims 1, 2 or 3, wherein said antibody is selected from the group consisting of: antibodies for treating cardiovascular disease, antibodies for treating

respiratory disease, antibodies for treating tissue transplant rejection, antibodies for treating organ transplant rejection, antibodies for treating cancer, antibodies for treating inflammatory disease and antibodies used in radioimmunotherapy.

17. The crystal according to any one of claims 1, 2 or 3, wherein said crystal is labelled.

18. The crystal according to claim 17, wherein said crystal is labelled with a label selected from the group consisting of radiolabels, enzyme labels, toxins, magnetic agents or drug conjugates.

19. A dried crystal of a whole antibody.

20. A dried crystal of a single-chain Fv fragment of an antibody or an Fab fragment of an antibody.

21. A composition for the release of a whole antibody, a single-chain Fv antibody fragment, or an Fab antibody fragment, said composition comprising:

(a) a whole antibody crystal, a single-chain Fv antibody fragment crystal, or an Fab antibody fragment crystal, and

(b) at least one polymeric carrier.

22. A formulation, said formulation comprising:

(a) a whole antibody crystal, a single-chain Fv antibody fragment crystal, or an Fab antibody

fragment crystal, and

(b) at least one ingredient.

23. A composition for the release of a whole antibody, a single-chain Fv antibody fragment, or an Fab antibody fragment, said composition comprising:

(a) a formulation, wherein said formulation comprises a whole antibody crystal, a single-chain Fv antibody fragment crystal, or an Fab antibody fragment crystal, and an ingredient; and

(b) at least one polymeric carrier.

24. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 1 mg/ml.

25. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 10.1 mg/ml.

26. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 20 mg/ml.

27. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 50 mg/ml.

28. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 100 mg/ml.

29. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 120 mg/ml.

30. The crystal according to any one of claims 1, 2 or 3, or the composition according to claim 21 or 23, or the formulation according to claim 22, wherein said crystal or composition or formulation has an antibody or antibody fragment crystal concentration greater than about 200 mg/ml.

31. The composition according to claim 21 or 23 or the formulation according to claim 22, wherein said antibody or antibody fragment is a therapeutic antibody or antibody fragment.



32. The composition according to claim 21 or 23, wherein said polymeric carrier is a biodegradable polymer.

33. The composition according to claim 21 or 23, wherein said polymeric carrier is a biocompatible polymer.

34. The composition according to claim 21 or 23, wherein said polymeric carrier is a polymer selected from one or more of the group consisting of: poly (acrylic acid), poly (cyanoacrylates), poly (amino acids), poly (anhydrides), poly (depsipeptide), poly (esters), poly (lactic acid), poly (lactic-co-glycolic acid) or PLGA, poly (b-hydroxybutyrate), poly (caprolactone), poly (dioxanone); poly (ethylene glycol), poly ((hydroxypropyl)methacrylamide, poly [(organo)phosphazene], poly (ortho esters), poly (vinyl alcohol), poly (vinylpyrrolidone), maleic anhydride-alkyl vinyl ether copolymers, pluronic polyols, albumin, alginate, cellulose and cellulose derivatives, collagen, fibrin, gelatin, hyaluronic acid, oligosaccharides, glycaminoglycans, sulfated polysaccharides, blends and copolymers thereof.

35. The composition according to claim 21 or 23, wherein said polymeric carrier is poly(lactic-co-glycolic acid).

36. The composition according to claim 21 or 23, wherein said polymeric carrier is emulsified with poly(vinyl alcohol).

37. The composition according to claim 21 or 23, wherein said polymeric carrier is a co-polymer.

38. The formulation according to claim 22 or the composition according to claim 23, wherein said ingredient is albumin.

39. The formulation according to claim 22 or the composition according to claim 23, wherein said ingredient is selected from the group consisting of sucrose, trehalose, lactitol, gelatin, hydroxypropyl- $\beta$ -cyclodextrin, methoxypolyethylene glycol and polyethylene glycol.

40. A method for treating a mammal comprising the step of administering to the mammal an effective amount of a whole antibody crystal, a single-chain Fv antibody fragment crystal, or an Fab antibody fragment crystal.

41. A method for treating a mammal comprising the step of administering to the mammal an effective amount of the composition according to claim 21 or 23, or the formulation according to claim 22.

42. The method according to claim 41, wherein the composition or formulation is administered by parenteral route, oral route, or by needle-free injection.

43. A large-batch crystallization method for crystallizing a whole antibody, a single-chain Fv

antibody fragment or an Fab antibody fragment, comprising the steps of:

(a) mixing a solution of a whole antibody, a single-chain Fv antibody fragment or an Fab antibody fragment with a crystallization solution or a crystallization buffer; and

(b) agitating said mixture for between about 3 and about 48 hours at a temperature between about -21°C and about 61°C, until crystals of said antibody or said antibody fragment are formed.

44. The large-batch crystallization method according to claim 43, further comprising the step of drying said crystals by a method selected from the group consisting of: air drying, spray drying, lyophilization, vacuum oven drying and nitrogen gas drying.

45. The large-batch crystallization method according to claim 43, wherein said temperature is between about 4°C and about 37°C.

46. The large-batch crystallization method according to claim 43, wherein said temperature is between about -20°C to about 4°C.

47. The large-batch crystallization method according to claim 43, wherein said temperature is between about 22°C and about 61°C.

48. The large-batch crystallization method according to claim 43, wherein the pH of said

crystallization buffer is within a range from about pH 1.9 to about pH 11.1.

49. The large-batch crystallization method according to claim 43, wherein the pH of said crystallization buffer is within a range from about pH 1.9 to about pH 4.0.

50. The large-batch crystallization method according to claim 43, wherein the pH of said crystallization buffer is between about pH 3 and about pH 10.

51. The large-batch crystallization method according to claim 43, wherein the pH of said crystallization buffer is within a range from about pH 9.0 to about pH 11.1.

52. The large-batch crystallization method according to claim 43, wherein the polyethylene glycol (PEG) concentration (w/v) between about 5 and about 40%.

53. The large-batch crystallization method according to claim 43, wherein said crystallization buffer contains a polyethylene glycol (PEG) concentration (w/v) between about 1.9% and about 80%.

54. The large-batch crystallization method according to claim 43, wherein said crystallization buffer contains a polyethylene glycol (PEG) concentration (w/v) between about 1.9% and about 5%.

55. The large-batch crystallization method according to claim 43, wherein said crystallization buffer contains a polyethylene glycol (PEG) concentration (w/v) between about 20% and about 81%.

56. The large-batch crystallization method according to claim 43, wherein said crystallization buffer comprises polyethylene glycol (PEG) of a size ranging between about 200 and about 20000.

57. The large-batch crystallization method according to claim 43, wherein said crystallization buffer comprises polyethylene glycol (PEG) of a size between about 200 and about 80,000.

58. The large-batch crystallization method according to claim 43, wherein said crystallization buffer comprises polyethylene glycol (PEG) of a size between about 200 to about 400.

59. The large-batch crystallization method according to claim 43, wherein said crystallization buffer comprises polyethylene glycol (PEG) of a size between about 400 to about 80,000.

60. The large-batch crystallization method according to claim 43, wherein the concentration in said solution of the antibody or single-chain Fv antibody fragment or Fab antibody fragment to be crystallized is between about 0.01 mg/ml and about 500 mg/ml.

61. The large-batch crystallization method according to claim 43, wherein the concentration of the antibody or single-chain Fv antibody fragment or Fab antibody fragment to be crystallized is between about 0.01 mg/ml and about 4 mg/ml.

62. The large-batch crystallization method according to claim 43, wherein the concentration of the antibody or single-chain Fv antibody fragment or Fab antibody fragment to be crystallized is between above about 10 mg/ml and about 25 mg/ml.

63. The large-batch crystallization method according to claim 43, wherein the concentration of the antibody or single-chain Fv antibody fragment or Fab antibody fragment to be crystallized is between about 3 mg/ml and about 200 mg/ml.

64. The large-batch crystallization method according to claim 43, wherein the concentration of the antibody or single-chain Fv antibody fragment or Fab antibody fragment to be crystallized is between above about 25 mg/ml and about 500 mg/ml.

65. The large-batch crystallization method according to claim 43, wherein said crystallization buffer has a salt content between about 10 mM and about 400 mM.

66. The large-batch crystallization method according to claim 43, wherein said crystallization buffer has a buffer concentration between about 0 mM

and about 4 M.

67. The large-batch crystallization method according to claim 43, wherein said crystallization buffer has a buffer concentration between about 0 mM and about 2 mM.

68. The large-batch crystallization method according to claim 43, wherein said crystallization buffer has a buffer concentration between about 1 M and about 4 M.

69. A method for purifying a protein by affinity matrix purification, said method comprising the steps of:

(a) mixing with a binding buffer crystals of a whole antibody, a single-chain Fv antibody fragment or an Fab antibody fragment, wherein said antibody or antibody fragment has affinity for the protein to be purified;

(b) adding a protein solution comprising the protein to be purified to the crystal/buffer mixture;

(c) incubating the protein/crystal/buffer mixture for a time and at a temperature sufficient to permit binding of the protein to the antibody or antibody fragment;

(d) washing the mixture with a wash buffer;  
and

(e) eluting the protein from the mixture with an elution buffer.

70. A diagnostic kit for the *in vitro*

detection of an antigen in a sample, said kit comprising:

(a) a crystal of a whole antibody, a crystal of a single-chain Fv antibody fragment or a crystal of an Fab antibody fragment, wherein said antibody fragment is capable of specifically binding to said antigen; and

(b) one or more reagents for detecting the binding of said antibody crystal or antibody fragment crystal to any antigen in said sample.

71. The diagnostic kit according to claim 70, wherein said antigen is a viral antigen.

72. An *in vitro* diagnostic method for detecting the presence of an antigen in a sample comprising the steps of:

(a) contacting said sample with a crystal of a whole antibody, a crystal of a single-chain Fv antibody fragment or a crystal of an Fab antibody fragment, wherein said antibody or antibody fragment is capable of specifically binding to said antigen, under conditions which permit said antibody crystal or antibody fragment crystal to bind to any antigen in said sample; and

(b) detecting the binding of said antibody crystal or antibody fragment to any antigen in said sample.

73. The diagnostic method according to claim 72, wherein said antigen is a viral antigen.



74. A large-batch crystallization method for crystallizing a whole antibody, a single-chain Fv antibody fragment or an Fab antibody fragment, comprising the steps of:

(a) mixing a solution of a whole antibody, a single-chain Fv antibody fragment or an Fab antibody fragment with a crystallization solution or crystallization buffer; and

(b) agitating said mixture for between about 5 minutes and about 72 hours at a temperature between about -21°C and about 61°C, until crystals of said antibody or said antibody fragment are formed.

75. The large-batch crystallization method according to claim 43, wherein said solution of antibody to be crystallized is produced by a method comprising the steps of:

(a) centrifuging transgenic milk comprising a whole antibody to remove milk fat and produce skim transgenic milk; and

(b) diluting the skim transgenic milk obtained in step (a) with about 250 mM EDTA to produce a solution of clarified skim transgenic milk comprising said antibody.

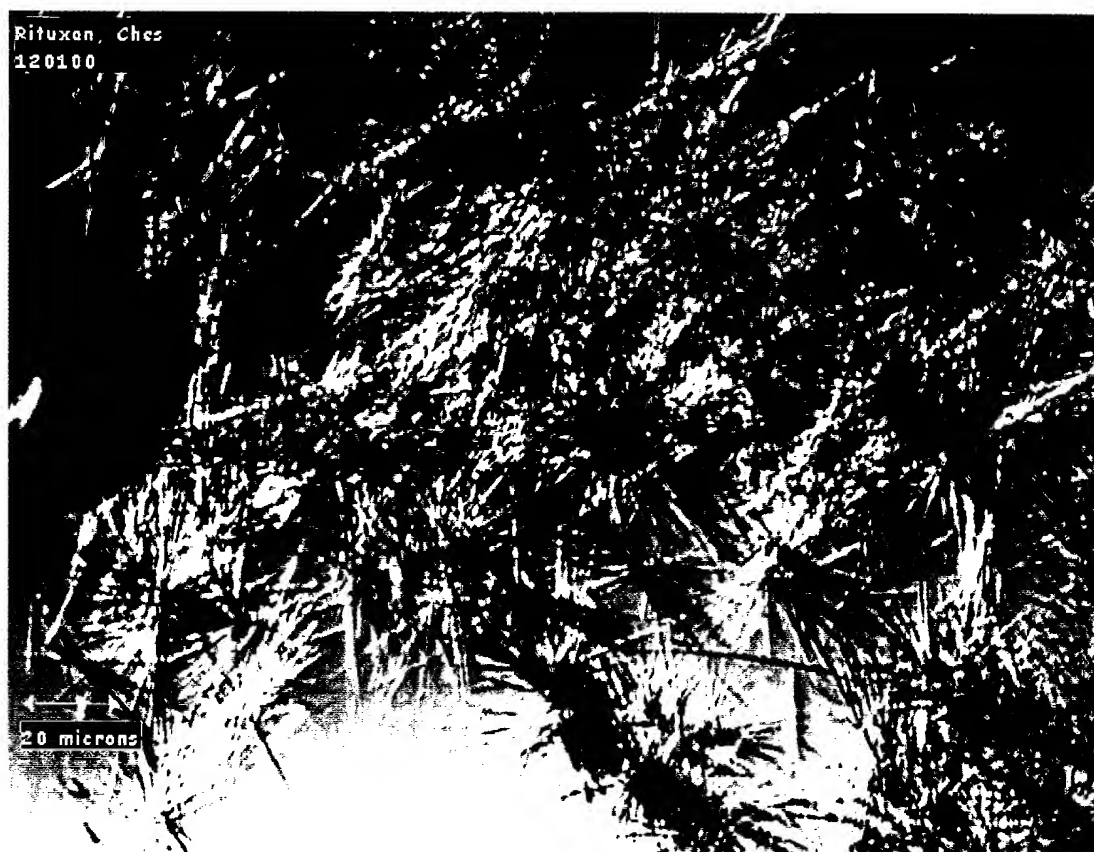
76. The composition according to claim 21 or 23, or the formulation according to claim 22, wherein said whole antibody crystal, single-chain Fv antibody fragment crystal, or Fab antibody fragment crystal is crosslinked.

77. A method for treating a mammal

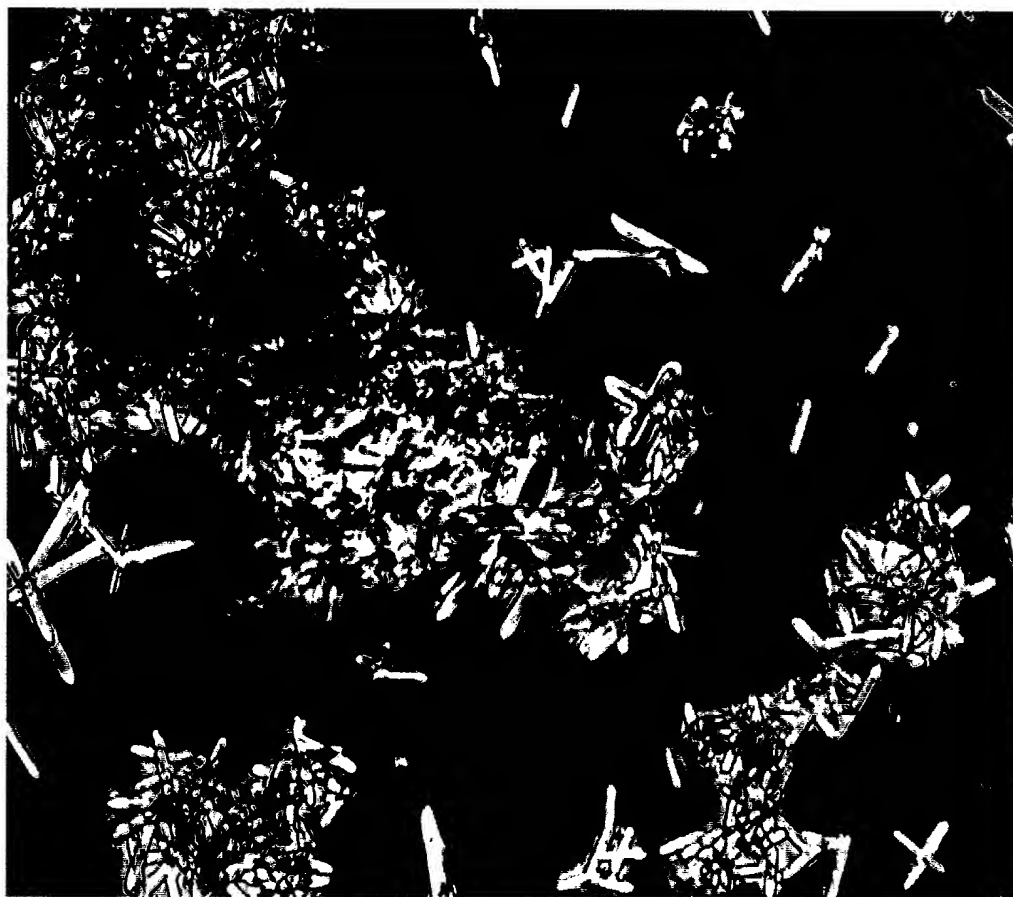
comprising the step of administering to the mammal an effective amount of the composition according to claim 76.

78. The method according to claim 77, wherein the composition is administered by parenteral route, oral route, or by needle-free injection.

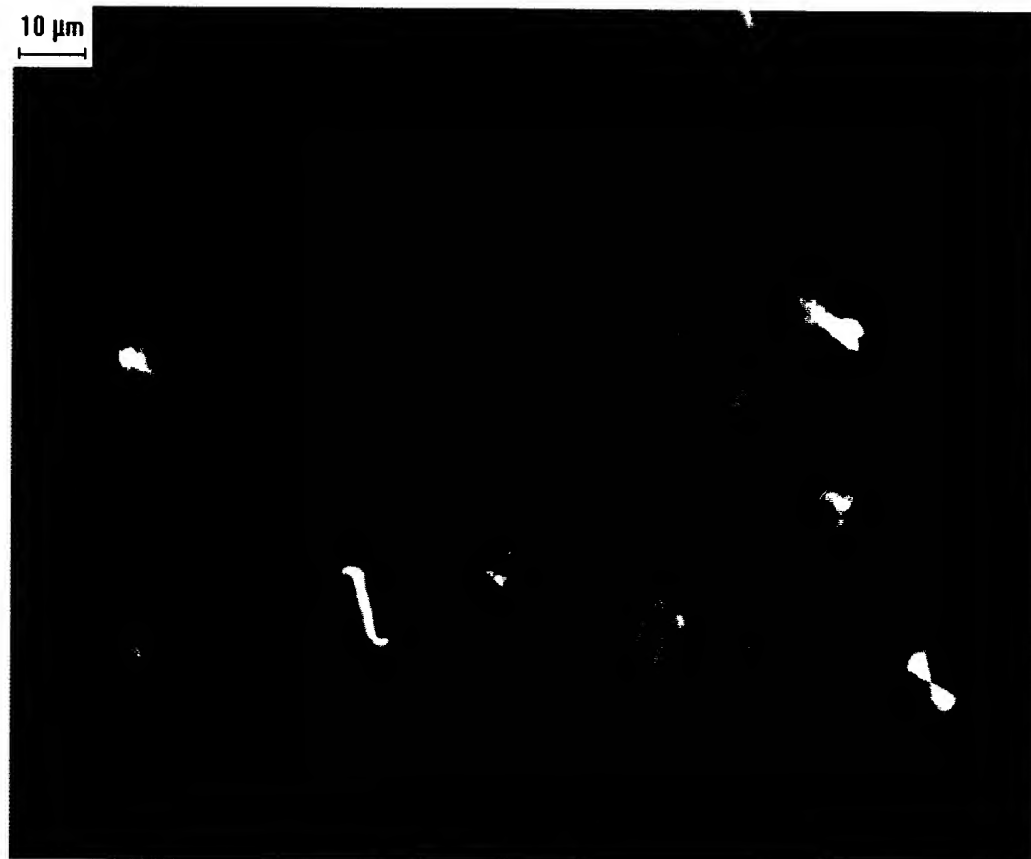
**Figure 1: Rituximab Crystals**



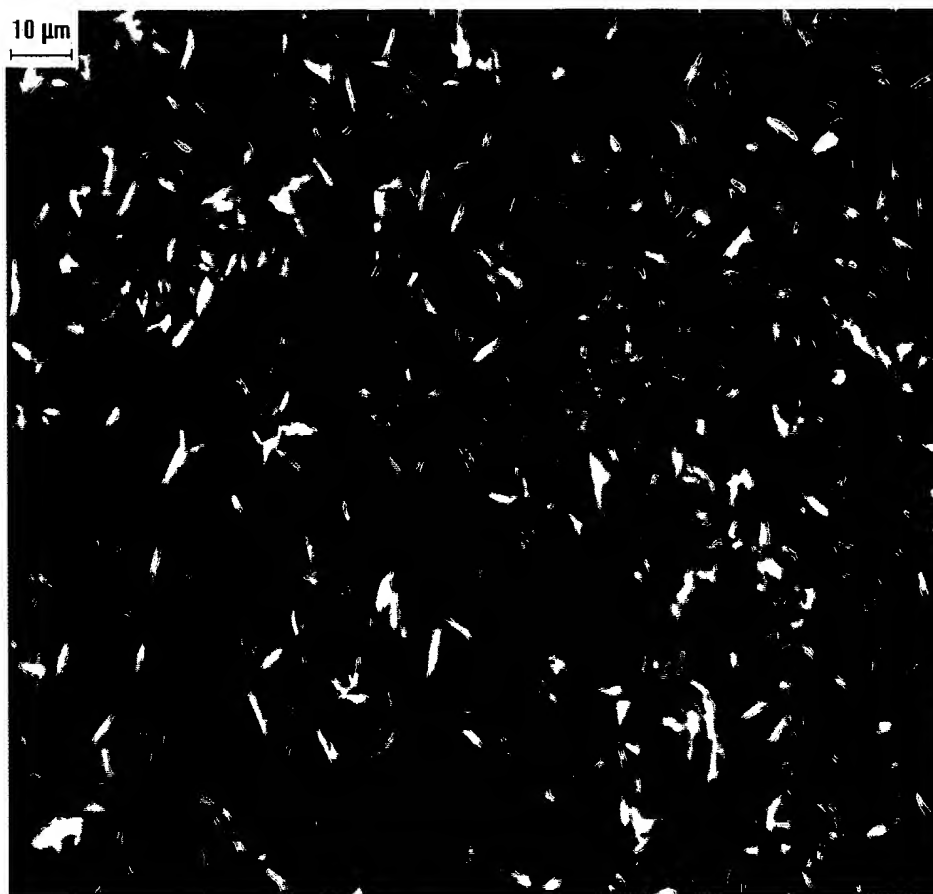
**Figure 2: Infliximab Crystals**



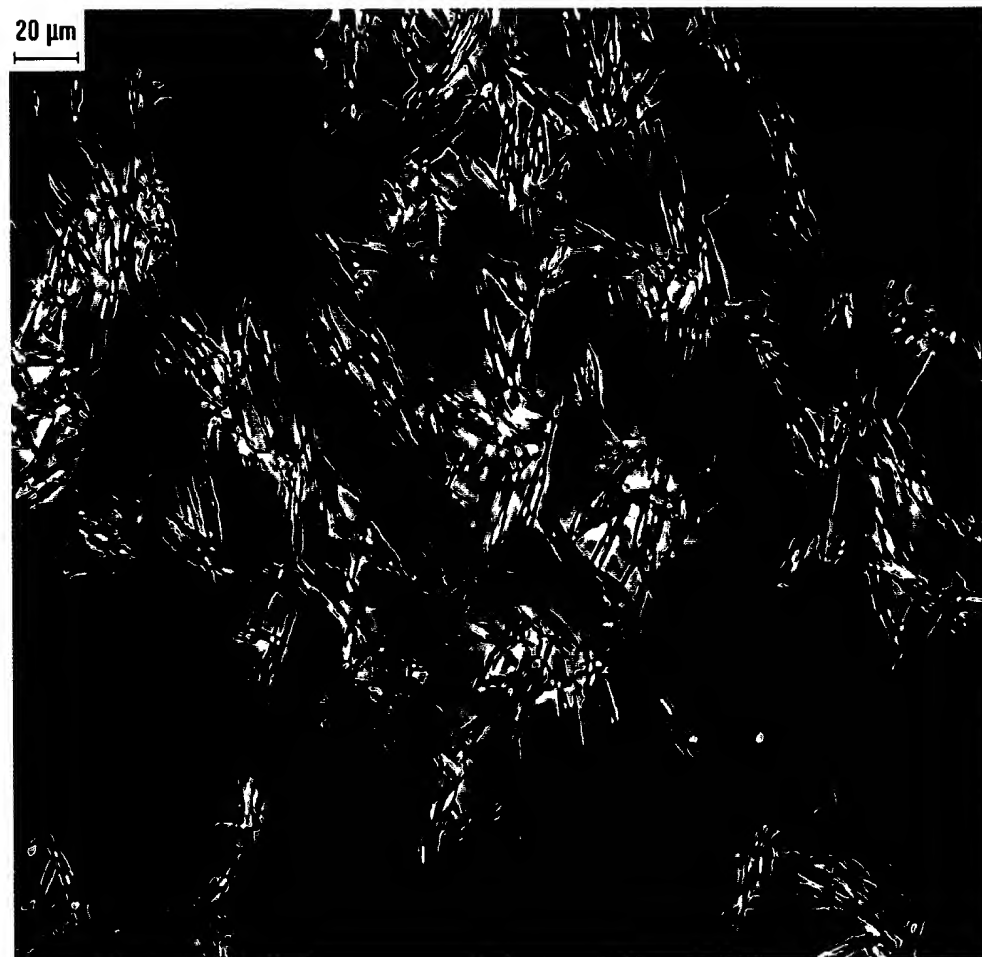
**Figure 3: Rituximab Cube-shaped Crystals**



**Figure 4: Rituximab Small Needle-like Crystals**



**Figure 5: Trastuzumab Crystals**

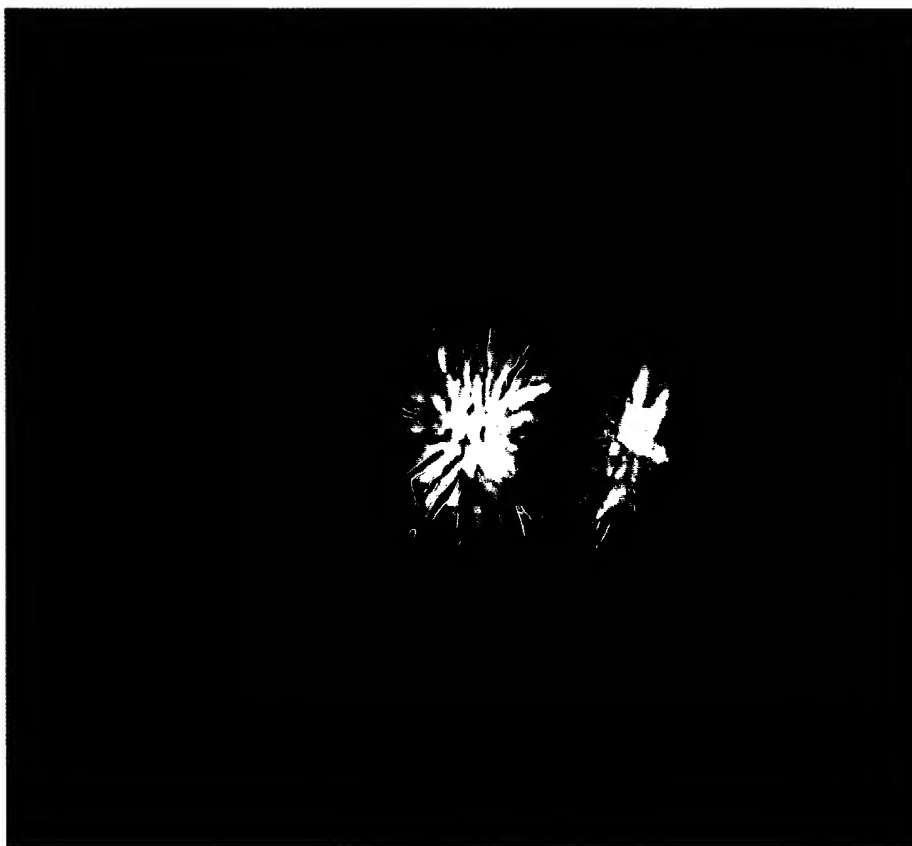


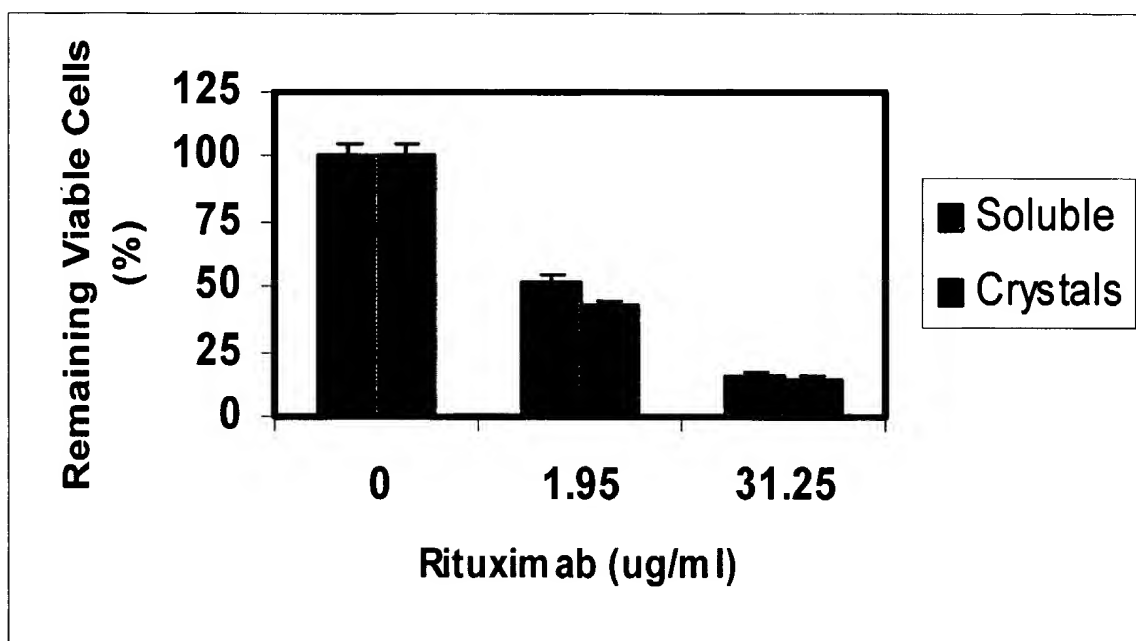
**Figure 6: Trastuzumab Long Needle-like Crystals**



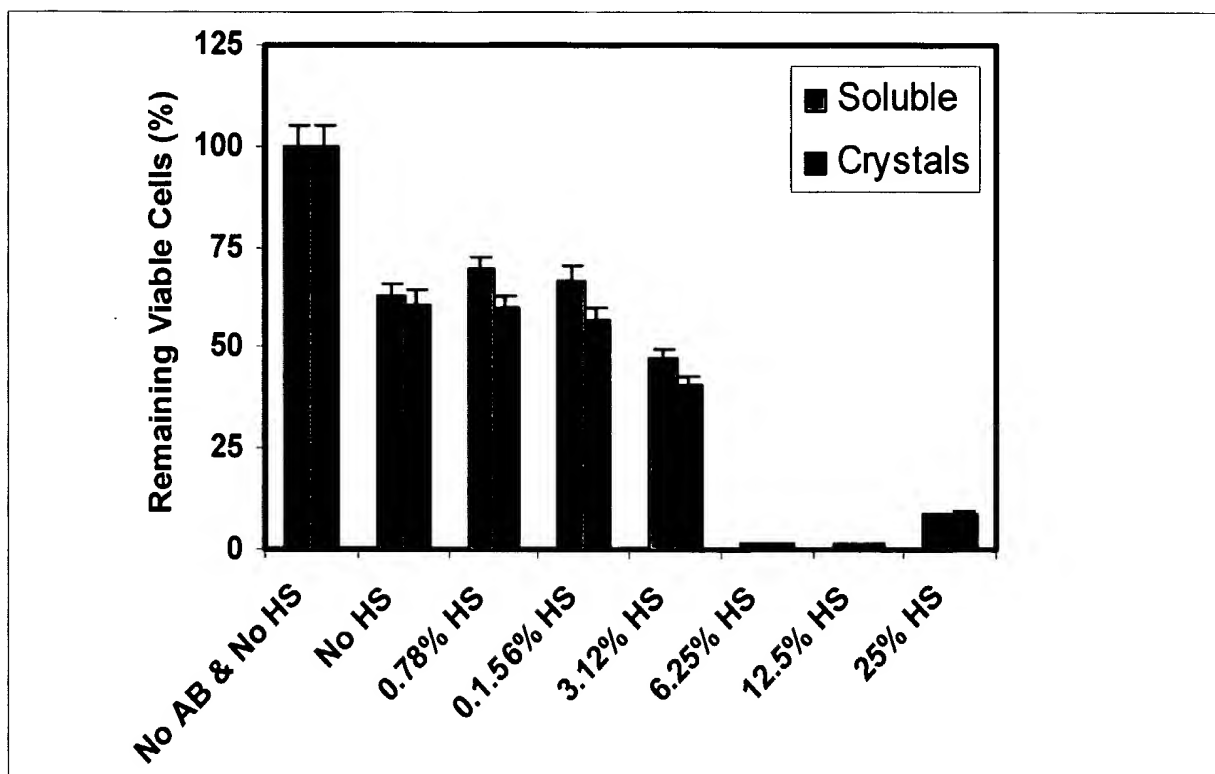


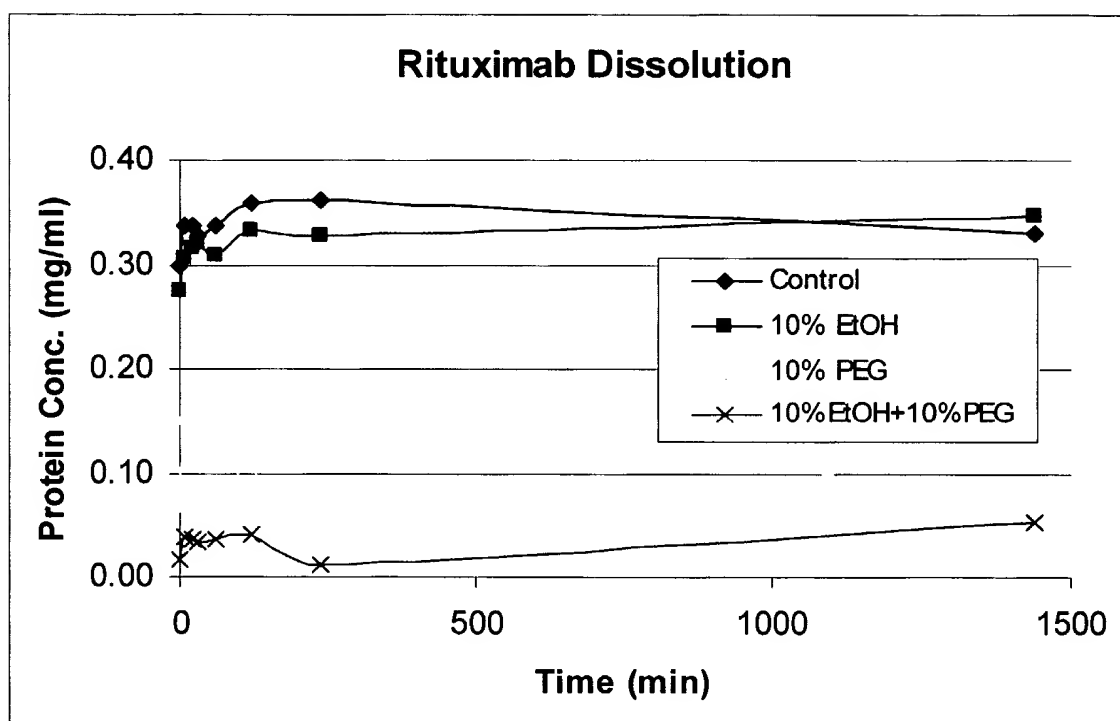
**Figure 7: Infliximab Star-shaped Crystals**

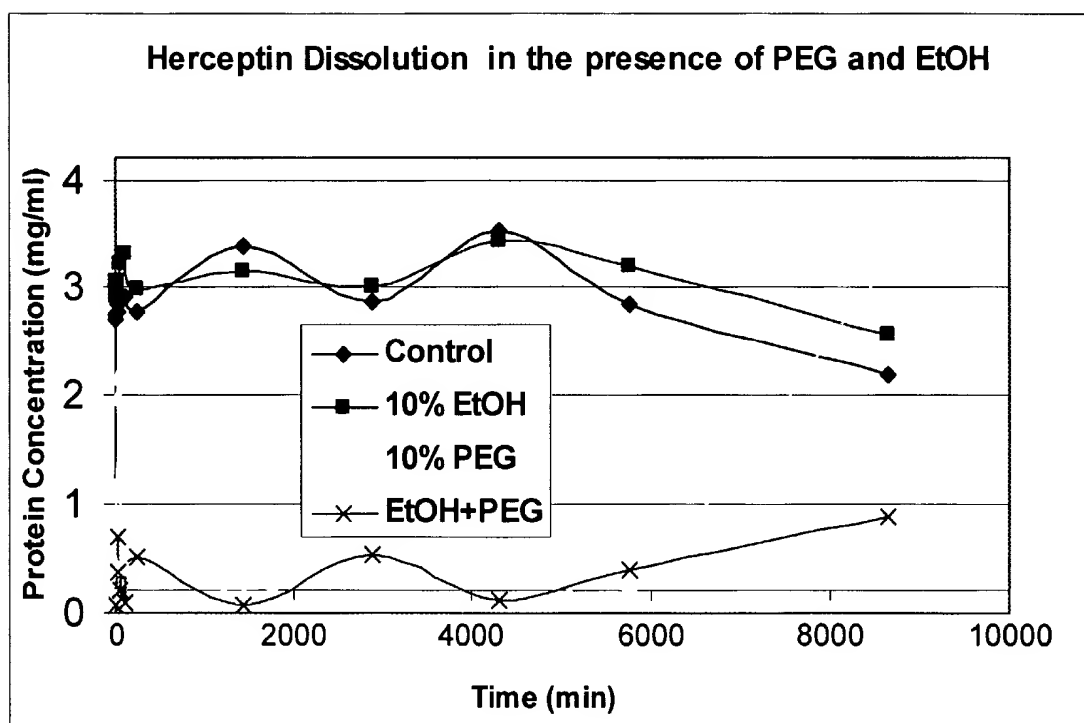


**Figure 8: Crystallized Rituximab-Induced Direct Cytotoxicity**

**Figure 9: Crystallized Rituximab-Induced Complement-Dependent Cytotoxicity**



**Figure 10: Analysis of Dissolved Rituximab Crystals**

**Figure 11: Analysis of Dissolved Trastuzumab Crystals**

**Figure 12: Stability of Rituximab in the Crystalline Form**

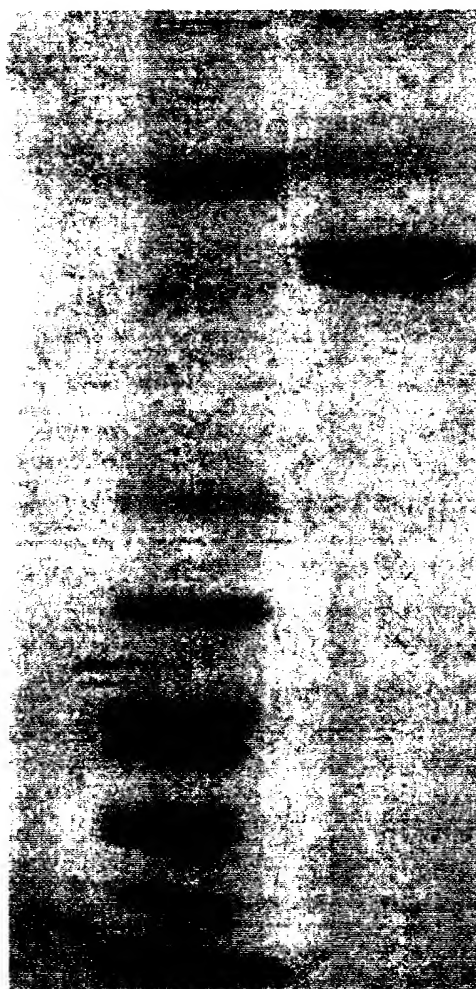


Figure 13: Crystalline Trastuzumab after treatment with Acetone

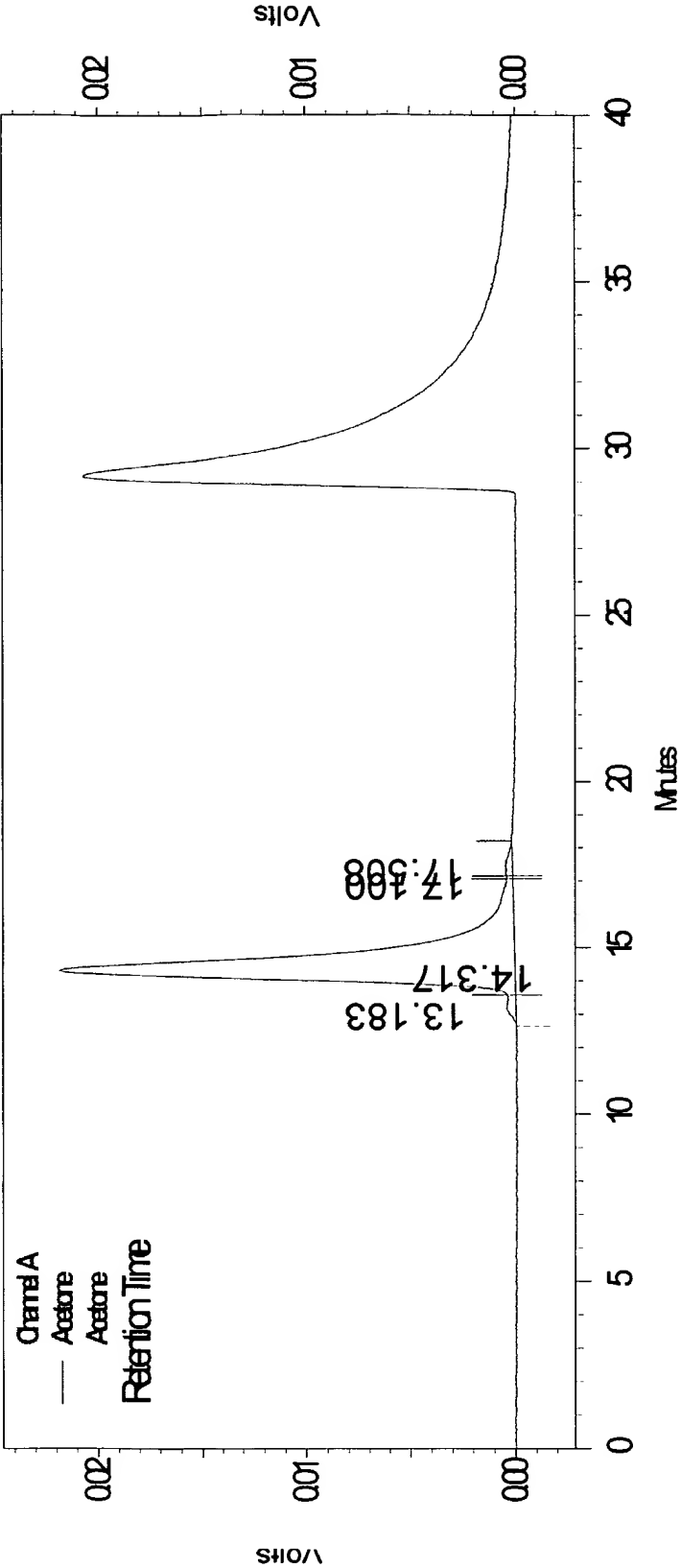
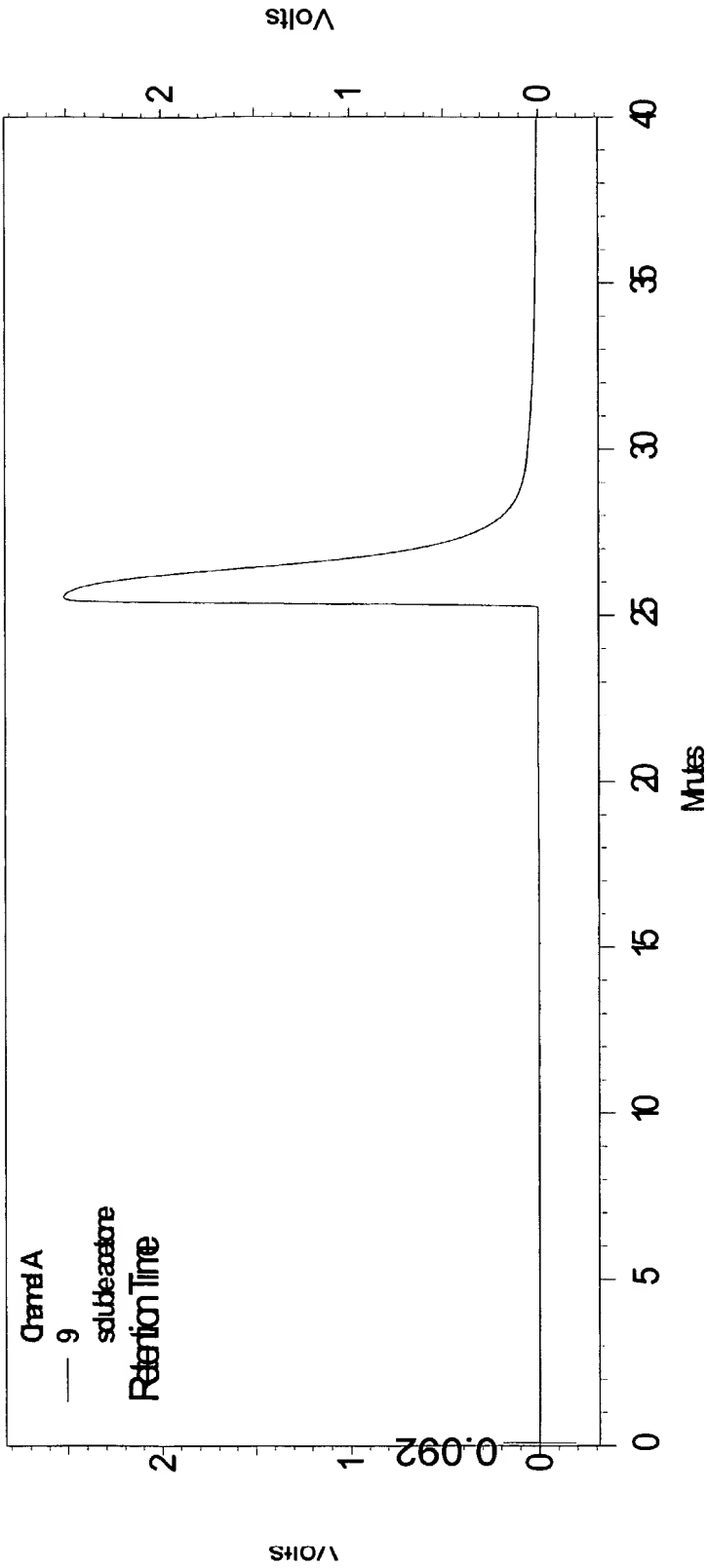
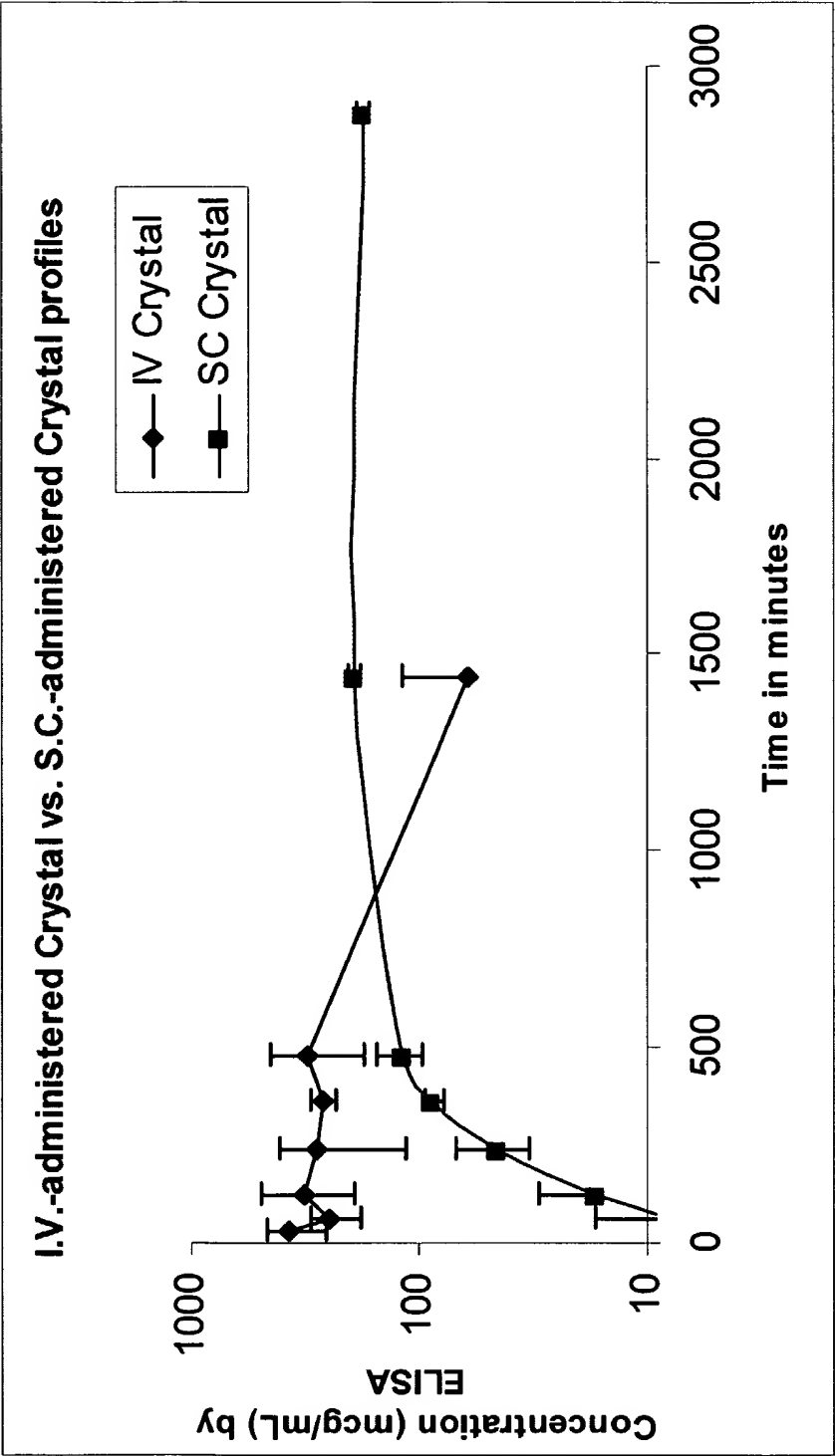


Figure 14: Soluble Trastuzumab after treatment with Acetone





**Figure 15: Bioavailability of Crystalline Trastuzumab Administered Intravenously (I.V.) or Subcutaneously (S.C.)**



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(81) Designated States (*national*): AE, AG, AL, AM, AT, AU,  
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[Continued on next page]

(54) Title: CRYSTALS OF WHOLE ANTIBODIES AND FRAGMENTS THEREOF AND METHODS FOR MAKING AND USING THEM

### Rituximab Crystals



(57) Abstract: Methods are also provided for preparing stabilized formulations of whole antibody crystals or antibody fragment crystals using pharmaceutical ingredients or excipients and optionally encapsulating the crystals or crystal formulations in a poly-  
meric carrier to produce compositions and using such protein crystals for biomedical applications, including delivery of therapeutic proteins and vaccines.



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## INTERNATIONAL SEARCH REPORT

International Application No

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## A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07K16/00 A61K39/395 G01N33/53 A61K47/06 A61K47/36  
 A61K47/38 A61K47/42 A61P35/00 G01N33/569

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

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Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

MEDLINE, EPO-Internal, WPI Data, PAJ, BIOSIS, CHEM ABS Data, EMBASE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category °	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>HARRIS L J ET AL: "Crystallographic structure of an intact IgG1 monoclonal antibody."            JOURNAL OF MOLECULAR BIOLOGY. ENGLAND 6            FEB 1998,            vol. 275, no. 5,            6 February 1998 (1998-02-06), pages            861-872, XP002208332            ISSN: 0022-2836            abstract            page 861, column 1, line 7 -column 2,            paragraph 2            page 869, column 1, paragraph 6 -column 2,            paragraph 1</p> <p style="text-align: center;">--- -/--</p>	1,4,11



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

° Special categories of cited documents:

\*A\* document defining the general state of the art which is not considered to be of particular relevance

\*E\* earlier document but published on or after the international filing date

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\*O\* document referring to an oral disclosure, use, exhibition or other means

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\*X\* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

\*Y\* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

\*&amp;\* document member of the same patent family

Date of the actual completion of the international search

19 November 2002

Date of mailing of the international search report

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Name and mailing address of the ISA

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## INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 01/49628

## C (Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>WO 99 55310 A (MARGOLIN ALEXEY L ;RAKESTRAW SCOTT L (US); KHALAF NAZER K (US); SH) 4 November 1999 (1999-11-04)</p> <p>abstract  page 1, line 4-12  page 1, line 25  page 2, line 3-32  page 3, line 26-33  page 8, line 20-33  page 9, line 10-29  page 10, line 1-28  page 14, line 4-27  page 15, line 11-16  page 16, line 3,4  page 16, line 27 -page 17, line 4  page 18, line 8-16  page 28, line 24 -page 29, line 13  page 29, line 14-18  page 30, line 18-26  page 36, line 22-29  page 37, line 16  page 39, line 31  page 41, line 1-12  page 51, line 22 -page 54, line 16  page 54, line 17-25  page 60, line 24 -page 61, line 24  page 66, line 26 -page 67, line 25  page 68, line 1 -page 69, line 7  page 71, line 1 -page 72, line 26  page 78, line 6 -page 79, line 5  page 81, line 25 -page 82, line 13  page 93, line 8 -page 94, line 16  page 112, line 15 -page 113, line 16  page 116, line 7 -page 118, line 18</p> <p>----</p>	<p>1,4,11, 21-30, 32-68, 74-78</p>
X	<p>US 6 011 001 A (NAVIA MANUEL A ET AL) 4 January 2000 (2000-01-04)</p>	<p>1,4,11, 17,18, 21-23, 33, 40-42, 70-73, 76-78</p>
Y	<p>abstract</p> <p>column 3, line 10-60  column 10, line 14-19  column 11, line 11-22  column 28, line 54 -column 29, line 23  column 32, line 10 -column 33, line 2  column 33, line 33-47  column 34, line 27-62  column 47, paragraphs 41-52  column 49, line 38-56  column 55, line 51-54</p> <p>----</p>	<p>43-68, 74,75</p>
	<p>-----</p> <p>-/--</p>	

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>US 6 140 475 A (MARGOLIN ALEXEY L ET AL) 31 October 2000 (2000-10-31)</p> <p>abstract column 6, line 42-61 column 8, line 19 -column 9, line 67 column 10, line 42 -column 11, line 18 column 15, line 1-10 column 32, line 46-62 column 34, line 65 -column 35, line 16 column 39, line 14-32</p> <p>---</p>	<p>1,4,11, 21-23, 32-34, 39-68, 74-78</p>
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X	<p>BRADEN B C ET AL: "X-ray crystal structure of an anti-Buckminsterfullerene antibody fab fragment: biomolecular recognition of C(60)." PROCEEDINGS OF THE NATIONAL ACADEMY OF SCIENCES OF THE UNITED STATES OF AMERICA. UNITED STATES 24 OCT 2000, vol. 97, no. 22, 24 October 2000 (2000-10-24), pages 12193-12197, XP002208334 ISSN: 0027-8424 abstract page 12193, column 2, paragraph 2</p> <p>---</p>	<p>3,4,11</p>
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	<p>---</p> <p>-/--</p>	

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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Y	US 6 042 824 A (KHALAF NAZER K) 28 March 2000 (2000-03-28) abstract column 5, line 24-51 column 6, line 1-13 column 7, line 28-66 column 9, line 10-49 column 11, line 47-51 column 12, line 14-16 column 13, line 5-21 column 13, line 67 -column 14, line 4	43-68, 74,75
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Y	WO 91 09943 A (NOVONORDISK AS) 11 July 1991 (1991-07-11) abstract page 2, line 1-30 page 4, line 17-23 page 5, line 15-23 page 10, line 14-16 page 11, line 1-15 page 12, line 15-19	43-68, 74,75
A	BLANCO DOLORES ET AL: "Protein encapsulation and release from poly(lactide-co-glycolide) microspheres: Effect of the protein and polymer properties and of the co-encapsulation of surfactants." EUROPEAN JOURNAL OF PHARMACEUTICS AND BIOPHARMACEUTICS, vol. 45, no. 3, May 1998 (1998-05), pages 285-294, XP002221361 ISSN: 0939-6411 abstract	1-4,11, 17,18, 21-30, 32-42, 70-73, 76-78

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US 01/49628

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:  
Although claims 40-42, 77 and 78 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the antibody crystals.
2. ☒ Claims Nos.: 5,7,16,31  
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:  
see FURTHER INFORMATION sheet PCT/ISA/210
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

see additional sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☒ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:  
1-4, 11, 17, 18, 21-30, 32-68, 70-78
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

### Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☒ No protest accompanied the payment of additional search fees.



**FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210**

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

Invention 1: claim 1 (complete), 4, 11, 17, 18,  
21-30, 32-42, 70-73 and 76-78 (all partial)

Crystals of a whole antibody; Compositions and formulations thereof; Methods for use.

Invention 2: claim 2 (complete), 4, 11, 17, 18,  
21-30, 32-42, 70-73 and 76-78 (all partial)

Crystals of single-chain Fv fragments of an antibody;  
Compositions and formulations thereof; Compositions and  
formulations thereof; Methods for use.

Invention 3: claim 3 (complete), 4, 11, 17, 18,  
21-30, 32-42, 70-73 and 76-78 (all partial)

Crystals of a Fab fragment of an antibody; Compositions and  
formulations thereof; Compositions and formulations thereof;  
Methods for use.

Inventions 4-48: claims 12, 19, 43-69, 74,  
75 (all complete), 4, 6, 8-11, 13-15, 17, 18,  
20-30, 32-42, 70-73, 76-78 (all partial)

Crystals of polyclonal, monoclonal, chimeric, humanised,  
non-glycosylated, bispecific, human, mouse, IgG, IgM, IgA,  
IgD, IgE, anti-idiotypic, Rituximab, Infliximab, Trastuzumab,  
Abciximab, Palivizumab, Murumonab-CD3, Gemtuzumab,  
Basiliximab, Daclizumab, Etanercept, Ibritumomab, anti-TNF,  
anti-CD3, anti-CD20, anti-CD25, anti-CD33, anti-CD40,  
anti-HER2, anti-HBV, anti-HAV, anti-HCV, anti-GPIIb/IIIa  
receptor, anti-RSV, anti-HIV, anti-HSV, anti-EBV antibodies;  
dried crystals of whole, single-chain Fv and Fab fragments.  
Compositions and formulations thereof; Methods for use. In  
addition, methods for large-batch crystallisation and for  
protein purification are considered to be separate  
inventions.

## FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.2

Claims Nos.: 5,7,16,31

Present claims 5, 7, 16 and 31 relate to a product defined by reference to a desirable characteristic or property, namely a crystal of a therapeutic antibody or an antibody used for the treatment of various diseases or the crystal is a carrier-free pharmaceutical controlled release crystal.

The claims cover all antibody crystals having this characteristic or property, whereas the application provides support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT for only a very limited number of such crystals. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible. Independent of the above reasoning, the claims also lack clarity (Article 6 PCT). An attempt is made to define the crystal by reference to a result to be achieved. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible. Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to the crystals of an whole antibody.

The applicant's attention is drawn to the fact that claims, or parts of claims, relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure.

## INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

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Information on patent family members

International Application No

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			US 5837513 A	17-11-1998

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**MARGOLIN, Alexey, L.** [US/US]; 193 Upland Avenue,  
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(74) Agents: **PIERRI, Margaret, A.** et al.; c/o Fish & Neave,  
1251 Avenue of the Americas, New York, NY 10020 (US).

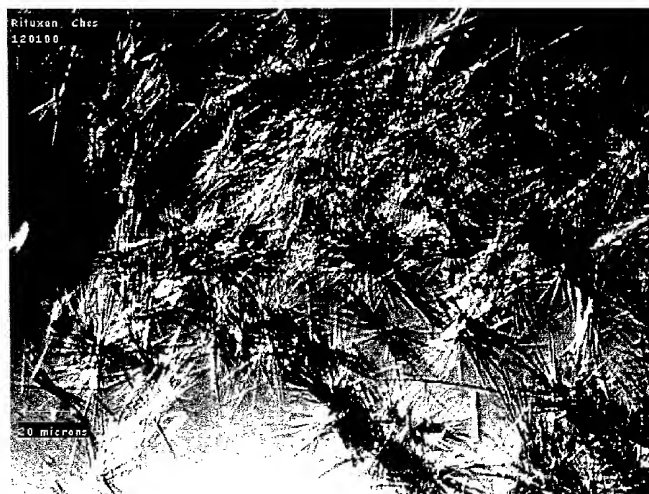
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AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU,  
CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,  
GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC,  
LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW,  
MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG,  
SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ,  
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KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW),  
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[Continued on next page]

(54) Title: CRYSTALS OF WHOLE ANTIBODIES AND FRAGMENTS THEREOF AND METHODS FOR MAKING AND USING THEM

#### Rituximab Crystals



(57) Abstract: Methods are also provided for preparing stabilized formulations of whole antibody crystals or antibody fragment crystals using pharmaceutical ingredients or excipients and optionally encapsulating the crystals or crystal formulations in a polymeric carrier to produce compositions and using such protein crystals for biomedical applications, including delivery of therapeutic proteins and vaccines.



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GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent  
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*For two-letter codes and other abbreviations, refer to the "Guid-  
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